

From: Snedden, Sheridan
Sent: Tuesday, August 19, 2003 6:48 PM
To: STIC-Biotech/ChemLib
Subject: Seq Search 09820053

Sheridan SNEDDEN ID# 79298 Date: 8/19 /2003

AU 1653

308-4843

Serial # : 09820053

Room Location: 10A12

Mail Box: 9B01

Claims reads: Peptide 5 to 23 amino acids length, 70% identical to SEQ ID NO: 43, wherein 80% of the amino acids are Phe, Lys, Leu and Ala.

I am not sure how to request this search. Please advise.

Thanks,
Examiner Snedden
#79298
A.U. 1653/ 9B01
Office Location: 10A12
Phone #: 305-4843

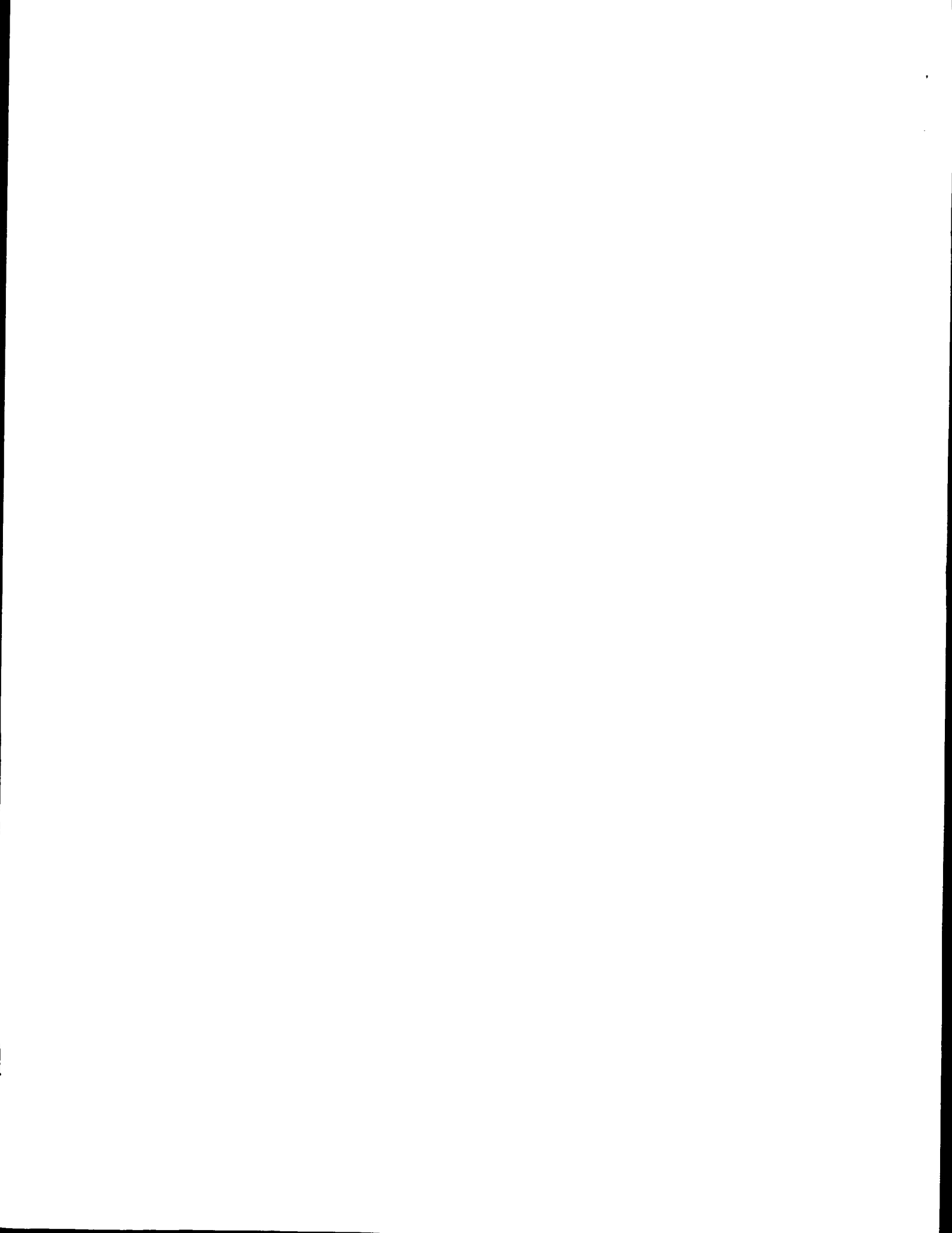
Searcher: D. S. L. 1653
Phone: 305-4843
Location: CM 1 6A03
Date Picked Up: _____
Date Completed: 8/22
Searcher Prep/Review: 15
Clerical: _____
Online time: 6

TYPE OF SEARCH:

NA Sequences: _____
AA Sequences: 1
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)

STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: Comp. Sys.
WWW/Internet: _____
Other (specify): _____





STIC Search Report

Biotech-Chem Library

STIC Database Tracking

TO: Sheridan Snedden
Location: CM1/10A12&9B01
Art Unit: 1653
Friday, August 22, 2003

Case Serial Number: 09/820053

From: David Schreiber
Location: Biotech-Chem Library
CM1-6A03
Phone: 308-4292

david.schreiber@uspto.gov

Search Notes

Sheridan,

Since this specifically refers to seq 43 and that sequence has the preferred residues, it seemed the best way to run this was just by doing a standard search of seq 43 and saving 45 alignments so you can see what you got. Hope this helps. Call if you have any questions.

David Schreiber
308-4292

5

207

59-50

60

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein protein search using sw model

Run on: August 21, 2004, 08:17:40 : Search time 28 seconds

(without alignments)
22.667 Million cell updates/sec

Title: us-09-820-053a-43

Perfect score: 66

Sequence: 1 FARALKALIKALKA15

Scoring table: RLSUM62

Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 4240858 residues

Total number of hits satisfying chosen parameters: 168489

Minimum DB seq length: 0
Maximum DB seq length: 25

Post processing: Minimum Match 100%
Maximum Match 100%
Listing first 45 summaries

Database :

IssuedPatents_AA: *
1: 2002-01-04 15:00:00 US-07-908-455A-6
2: 2002-01-04 15:00:00 US-07-908-455A-6
3: 2002-01-04 15:00:00 US-07-908-455A-6
4: 2002-01-04 15:00:00 US-07-908-455A-6
5: 2002-01-04 15:00:00 US-07-908-455A-6
6: 2002-01-04 15:00:00 US-07-908-455A-6

prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	51	77.3	21	1 US-07-908-455A-6	Sequence 6, Appl
2	51	77.3	21	1 US-08-434-120-32	Sequence 32, Appl
3	51	77.3	21	1 US-08-465-325-32	Sequence 32, Appl
4	51	77.3	21	4 US-09-115-737-32	Sequence 32, Appl
5	50	75.8	14	1 US-08-944-133-16	Sequence 16, Appl
6	50	75.8	14	1 US-08-944-133-33	Sequence 33, Appl
7	50	75.8	21	1 US-08-944-133-17	Sequence 17, Appl
8	50	75.8	21	1 US-08-944-133-30	Sequence 30, Appl
9	50	75.8	21	1 US-08-944-133-34	Sequence 34, Appl
10	48	72.7	23	1 US-08-231-730A-39	Sequence 39, Appl
11	48	72.7	23	1 US-08-427-001C-44	Sequence 44, Appl
12	48	72.7	23	1 US-08-457-171-39	Sequence 39, Appl
13	48	72.7	23	2 US-08-723-306-19	Sequence 19, Appl
14	48	72.7	23	2 US-08-505-486-44	Sequence 44, Appl
15	48	72.7	23	3 US-08-689-489C-39	Sequence 39, Appl
16	48	72.7	23	3 US-08-801-028-44	Sequence 44, Appl
17	48	72.7	23	3 US-09-340-154-44	Sequence 44, Appl
18	48	72.7	23	4 US-09-242-902A-39	Sequence 39, Appl
19	48	72.7	23	4 US-09-482-611B-44	Sequence 44, Appl
20	48	72.7	23	4 US-09-019-922A-44	Sequence 44, Appl
21	48	72.7	23	5 PCT-US94-12550-47	Sequence 47, Appl
22	48	72.7	23	5 PCT-US95-04335-39	Sequence 39, Appl
23	48	72.7	23	5 PCT-US95-04718-39	Sequence 39, Appl
24	48	72.7	23	5 PCT-US95-09338-44	Sequence 44, Appl
25	48	72.7	23	5 PCT-US95-09339-44	Sequence 44, Appl
26	48	72.7	23	5 PCT-US96-10041-19	Sequence 19, Appl
27	40	60.6	21	1 US-07-908-455A-6	Sequence 8, Appl

28 40 60.6 21 1 US-08-444 120-34
29 40 60.6 21 1 US-08-465-425-34
30 40 60.6 21 4 US-09-115 737-32
31 39 59.1 22 4 US-08-752 526-3
32 39 59.1 23 1 US-08-241 750A-42
33 39 59.1 23 2 US-08-505 486-47
34 39 59.1 23 3 US-08-689 489C-39
35 39 59.1 23 4 US-08-801 028-44
36 39 59.1 23 3 US-09-340 154-44
37 39 59.1 23 4 US-09-242 902A-39
38 39 59.1 23 4 US-09-482 611B-44
39 39 59.1 23 5 PCT-US95 09338-44
40 39 59.1 23 5 PCT-US95 09339-44
41 39 59.1 23 5 PCT-US96 10041-19
42 38 57.6 14 1 US-08-231-730A-39
43 48 57.6 21 1 US-08-444 120-34
44 48 57.6 21 1 US-07-908-455A-6
45 38 57.6 21 1 US-08-444 120-34

ALIGNMENTS

RESULT 1
US-07-908-455A-6
Sequence 6, Application US-07908455A

Patent No. 5459237

GENERAL INFORMATION:

APPLICANT: Beckwith, Barry A.

APPLICANT: Kari, U. Prasad

APPLICANT: Mallow, W. Lee

TITLE OF INVENTION: Peptide Compositions and

TITLE OF INVENTION: Uses Thereof

NUMBER OF SEQUENCES: 89

CORRESPONDENCE ADDRESS:

ADDRESSEE: Carella, Byrne, Rains, Williams,

ADDRESSEE: Carelli & Stewart

STREET: 6 Becker Farm Road

CITY: Roseland

STATE: New Jersey

COUNTRY: USA

ZIP: 07068

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch diskette

COMPUTER: IBM PS/2

OPERATING SYSTEM: PC-DOS

SOFTWARE: USA V2

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US-07-908-455A

FILING DATE: 19920702

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07686115

FILING DATE: 15-APR-1991

APPLICATION NUMBER: US 07476629

FILING DATE: 08-FEB-1990

ATTORNEY/AGENT INFORMATION:

NAME: Gistino, Elliot M.

REGISTRATION NUMBER: 24,025

REFERENCE/DEPOSIT NUMBER: 421250-122

TELECOMMUNICATION INFORMATION:

TELEPHONE: 201-994-1700

TELEFAX: 201-994-1744

INFORMATION FOR SEQ ID NO: 6:

SEQUENCE CHARACTERISTICS:

LENGTH: 21 amino acids

TYPE: AMINO ACID

STANDARDNESS:

TOPOLAGY: Linear

MOLECULE TYPE: Peptide

FEATURE:

OTHER INFORMATION: amide-terminated

US-07-908-455A-6

Query Match: 77.8% Score 51; DB 1; Length 21;
 Best Local Similarity: 85.7%; Pref. Rs. 0.2;
 Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 AKALKALKALKAL 16

DB 4 SEAKALKALKAL 17

RESULT 2

US 08 444 120 42

Sequence 42, Application US/08444120

Patent No. 5665479

GENERAL INFORMATION:

APPLICANT: Baker, Margaret A.

APPLICANT: Jacob, Leonard S.

APPLICANT: Maloy, W. Lee

TITLE OF INVENTION: Treatment of Gynecological

TITLE OF INVENTION: Malignancies with

TITLE OF INVENTION: Biologically Active Peptides

NUMBER OF SEQUENCES: 117

CORRESPONDENCE ADDRESS:

ADDRESSEE: Carolina, Ryngaert, Gilliland,

ADDRESSEE: Cecchi & Stewart

STREET: 6 Becker Farm Road

CITY: Roseland

STATE: New Jersey

COUNTRY: USA

ZIP: 07068

COMPUTER REARABLE FORM:

MEDIUM TYPE: 3.5 inch diskette

COMPUTER: IBM PS/2

OPERATING SYSTEM: PC-DOS

SOFTWARE: DW4.V2

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/92/444,120

FILING DATE:

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/297,950

FILING DATE:

APPLICATION NUMBER: US/98/226,108

FILING DATE:

APPLICATION NUMBER: US/97/947,462

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Olstein, Elliot M.

REGISTRATION NUMBER: 24,026

REFERENCE/AGENT NUMBER: 131,000,194

TELECOMMUNICATION INFORMATION:

TELEPHONE: 201-954-1700

TELEFAX: 201-994-1744

INFORMATION FOR SEQ ID NO: 42:

SEQUENCE CHARACTERISTICS:

LENGTH: 21 amino acids

TYPE: amino acid

STRANDINESS:

TOPOLOGY: linear

MOLECULE TYPE: peptide

US 08 444 120 42

Query Match: 77.8% Score 51; DB 1; Length 21;
 Best Local Similarity: 85.7%; Pref. Rs. 0.2;
 Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 AKALKALKALKAL 16

DB 4 SEAKALKALKAL 17

RESULT 3

US-08-465 425 42

Sequence 42, Application US/08465425

Patent No. 5665563

GENERAL INFORMATION:

APPLICANT: Marinin Pharmaceuticals Inc.

APPLICANT: 5110 Campus Drive

APPLICANT: Plymouth Meeting, PA 19462

TITLE OF INVENTION: Biologically Active Peptides

TITLE OF INVENTION: Biologically Active Peptides

NUMBER OF SEQUENCES: 114

CORRESPONDENCE ADDRESS:

ADDRESSEE: Finomson, Henderson, Paradowski

ADDRESSEE: Finomson

STREET: 1400 L Street, N.W. Suite 200

CITY: Washington

STATE: D.C.

COUNTRY: USA

ZIP: 20005 4415

MEDIUM TYPE: Floppy disk

COMPUTER: IBM compatible

OPERATING SYSTEM: PC-DOS

SOFTWARE: Patent in Release #1.0, Version #1.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/99/465,425

FILING DATE: 05 JUN 1999

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/184,412

FILING DATE: 18 JAN 94

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/994,201

FILING DATE: 01 JUN 92

ATTORNEY/AGENT INFORMATION:

NAME: Fordin, John W.

REGISTRATION NUMBER: 42,994

REFERENCE/AGENT NUMBER: 05,095,001

TELECOMMUNICATION INFORMATION:

TELEPHONE: (202) 408 4000

TELEFAX: (202) 408 4400

INFORMATION FOR SEQ ID NO: 42:

SEQUENCE CHARACTERISTICS:

LENGTH: 21 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: peptide

US 08 465 425 42

Query Match:

Best Local Similarity: 85.7%; Pref. Rs. 0.2;

Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 AKALKALKALKAL 16

DB 4 SEAKALKALKAL 17

RESULT 4

US 09 115 747 42

Sequence 42, Application US/09115742

Patent No. 6,646,445

GENERAL INFORMATION:

APPLICANT: D. Prasad Kati

APPLICANT: L. J. Williams

APPLICANT: Michael Melano

TITLE OF INVENTION: Biologically Active Peptides

TITLE OF INVENTION: Biologically Active Peptides

NUMBER OF SEQUENCES: 156

CORRESPONDENCE ADDRESS:

ADDRESSEE: Finomson, Henderson, Paradowski

ADDRESSEE: Finomson

STREET: 1400 L Street, N.W. Suite 200

CITY: Washington

STATE: D.C.

us-09-820-053a-43.ra1

Thu Aug 21 08:36:29 2003

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1  COUNTRY: USA
2  ZIP: 20005-3315
3  COMPUTER READABLE FORM:
4  MEDIUM TYPE: Floppy disk
5  OPERATING SYSTEM: PC-DOS/MS-DOS
6  SOFTWARE: Patulin Release
7  CURRENT APPLICATION NUMBER: US/09/115,737
8  FILING DATE: 15-JUL-1998
9  CLASSIFICATION: <Unknown>
10 PRIOR APPLICATION DATA:
11 APPLICATION NUMBER: 03/465,330
12 FILING DATE: 05-JUN-1995
13 APPLICATION NUMBER: 08/184,462
14 FILING DATE: 18-JAN-94
15 APPLICATION NUMBER: 07/891,201
16 FILING DATE: 01-JUN-92
17 ATTORNEY/AGENT INFORMATION:
18 NAME: Fordis, Jean B
19 REGISTRATION NUMBER: 32,984
20 REFERENCE/SEQUENCE NUMBER: 05387 0021-06000
21 TELECOMMUNICATION INFORMATION:
22 TELEPHONE: (202) 408-4000
23 TELEFAX: (202) 408-4400
24 INFORMATION FOR SEQ ID NO: 32:
25 SEQUENCE CHARACTERISTICS:
26 LENGTH: 21 amino acids
27 TYPE: amino acid
28 TOPOLOGY: linear
29 MOLECULE TYPE: peptide
30 SEQUENCE DESCRIPTION: SEQ ID NO: 32:
31
32 US-09-115-737-42
33
34 Query Match: 77.3%; Score 51; DB 4; Length 21;
35 Best Local Similarity: 85.7%; Pred. No. 0.2;
36 Matches: 12; Conservative: 1; Mismatches: 1; Indels: 1; Gaps: 0;
37
38 QY 2 AKALKALKALKAL 15
39 DB 4 SKALKALKALKAL 17
40
41 RESULT 5
42 US-08-944-133-43
43 Sequence 16: Application US/08944133
44 Patent No. 5789542
45 GENERAL INFORMATION:
46 APPLICANT: McLaughlin, Mark L
47 APPLICANT: Becker, Calvin L
48 TITLE OF INVENTION: Amphipathic Peptides
49 NUMBER OF SEQUENCES: 54
50 CORRESPONDENCE ADDRESS:
51 ADDRESSEE: John H. Rannels
52 STREET: P. O. Box 2471
53 CITY: Baton Rouge
54 STATE: LA
55 COUNTRY: USA
56 ZIP: 70821-2471
57 COMPUTER READABLE FORM:
58 MEDIUM TYPE: Floppy disk
59 COMPUTER: IBM PC compatible
60 OPERATING SYSTEM: PC-DOS/MS-DOS
61 SOFTWARE: Patent Release #1.0, Version #1.1.1.1
62 CURRENT APPLICATION DATA:
63 APPLICATION NUMBER: US/08/944,133
64 FILING DATE: 06-OCT-1997
65 CLASSIFICATION: 5540
66 PRIOR APPLICATION DATA:
67 APPLICATION NUMBER: 08/789,077
68 FILING DATE: 03-FEB-1997
69 APPLICATION NUMBER: 05/08/681,075
70 FILING DATE:
71 APPLICATION NUMBER: US/08/232,545
72 FILING DATE: 22-APR-1994
73 ATTORNEY/AGENT INFORMATION:
74 NAME: Rannels, John H
75 REGISTRATION NUMBER: 34451
76 REFERENCE/SEQUENCE NUMBER: ATTY File No. 5789542 9401
77 TELECOMMUNICATION INFORMATION:
78 TELEPHONE: 504 387-4221
79 TELEFAX: 504 346-8049
80 INFORMATION FOR SEQ ID NO: 33:
81 SEQUENCE CHARACTERISTICS:
82 LENGTH: 14 amino acids
83 TYPE: amino acid
84 STRANDEDNESS: Single
85 TOPOLOGY: linear
86 MOLECULE TYPE: peptide
87
88 US-08-944-133-16
89 Query Match: 75.8%; Score 50; DB 1; Length 14;
90 Best Local Similarity: 92.8%; Pred. No. 0.19;
91 Matches: 12; Conservative: 0; Mismatches: 1; Indels: 0; Gaps: 0;
92
93 QY 3 KALKALKALKAL 15
94 DB 1 KALKALKALKAL 13
95
96 RESULT 6
97 US-08-944-133-43
98 Sequence 43: Application US/08944133
99 Patent No. 5789542
100 GENERAL INFORMATION:
101 APPLICANT: McLaughlin, Mark L
102 APPLICANT: Becker, Calvin L
103 TITLE OF INVENTION: Amphipathic Peptides
104 NUMBER OF SEQUENCES: 54
105 CORRESPONDENCE ADDRESS:
106 ADDRESSEE: John H. Rannels
107 STREET: P. O. Box 2471
108 CITY: Baton Rouge
109 STATE: LA
110 COUNTRY: USA
111 ZIP: 70821-2471
112 COMPUTER READABLE FORM:
113 MEDIUM TYPE: Floppy disk
114 COMPUTER: IBM PC compatible
115 OPERATING SYSTEM: PC-DOS/MS-DOS
116 SOFTWARE: Patent Release #1.0, Version #1.1.1.1
117 CURRENT APPLICATION DATA:
118 APPLICATION NUMBER: US/08/944,133
119 FILING DATE: 06-OCT-1997
120 CLASSIFICATION: 5540
121 PRIOR APPLICATION DATA:
122 APPLICATION NUMBER: 08/789,077
123 FILING DATE: 03-FEB-1997
124 APPLICATION NUMBER: 05/08/681,075
125 FILING DATE:
126 APPLICATION NUMBER: US/08/232,545
127 FILING DATE: 22-APR-1994
128 ATTORNEY/AGENT INFORMATION:
129 NAME: Rannels, John H
130 REGISTRATION NUMBER: 34451
131 REFERENCE/SEQUENCE NUMBER: ATTY File No. 5789542 9401
132 TELECOMMUNICATION INFORMATION:
133 TELEPHONE: 504 387-4221
134 TELEFAX: 504 346-8049
135 INFORMATION FOR SEQ ID NO: 34:
136 SEQUENCE CHARACTERISTICS:
137 LENGTH: 14 amino acids
138 TYPE: amino acid
139 STRANDEDNESS: Single
140 TOPOLOGY: linear
141 MOLECULE TYPE: peptide

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US-08-944-133 43

Query Match 75.8%; Score 50; DB 1; Length 14;
 Best Local Similarity 92.4%; Prod. No. 0.19;
 Matches 12; Conservative 0; Mismatches 1; Indels 0;

QY 4 KALKALKALKAL 16
 11111 11111
 DB 2 KALKALKALKAL 14

RESULT 7

US-08-944-133 17
 Sequence 17, Application US/08944133

Patent No. 5789542

GENERAL INFORMATION:

APPLICANT: McLaughlin, Mark L.
 TITLE OF INVENTION: Amphipathic Peptides
 NUMBER OF SEQUENCES: 54

CORRESPONDENCE ADDRESS:

ADDRESSEE: John H. Rummels

STREET: P. O. Box 2471

CITY: Baton Rouge

STATE: LA

COUNTRY: USA

ZIP: 70821 2471

COMPUTER PEAKABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: IBM PC compatible

SOFTWARE: Patent in Release #1.0, Version #1.26

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/38/344,133

FILING DATE: 06 OCT 1997

CLASSIFICATION: 5540

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/789,077

FILING DATE: 03 FEB 1997

APPLICATION NUMBER: 02/38/0681,076

FILING DATE:

APPLICATION NUMBER: US/38/242,525

FILING DATE: 22 APR 1994

ATTORNEY/AGENT INFORMATION:

NAME: Rummels, John H

REGISTRATION NUMBER: 33451

REFERENCE/PACKET NUMBER: Atty File No. 5789542 9301

TELECOMMUNICATION INFORMATION:

TELEPHONE: 504 387-3221

TELEFAX: 504 346-8049

INFORMATION FOR SEQ ID NO: 17:

SEQUENCE CHARACTERISTICS:

LENGTH: 21 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-08-944-133 17

Query Match 75.8%; Score 50; DB 1; Length 21;
 Best Local Similarity 92.3%; Prod. No. 0.28;
 Matches 12; Conservative 0; Mismatches 1; Indels 0;

QY 4 KALKALKALKAL 15
 11111 11111
 DB 1 KALKALKALKAL 13

RESULT 8

US-08-944-133 40

Sequence 30, Application US/08944133

Patent No. 5789542

GENERAL INFORMATION:

APPLICANT: McLaughlin, Mark L.
 TITLE OF INVENTION: Amphipathic Peptides
 NUMBER OF SEQUENCES: 54
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: John H. Rummels
 STREET: P. O. Box 2471
 CITY: Baton Rouge
 STATE: LA
 COUNTRY: USA
 ZIP: 70821 2471
 COMPUTER PEAKABLE FORM:
 MEDIUM TYPE: Floppy disk
 OPERATING SYSTEM: IBM PC compatible
 SOFTWARE: Patent in Release #1.0, Version #1.26

CURRENT APPLICATION DATA:

APPLICATION NUMBER: 08/789,077

FILING DATE: 03 FEB 1997

APPLICATION NUMBER: US/38/344,133

FILING DATE:

APPLICATION NUMBER: 02/38/0681,076

FILING DATE:

APPLICATION NUMBER: US/38/242,525

FILING DATE: 22 APR 1994

ATTORNEY/AGENT INFORMATION:

NAME: Rummels, John H

REGISTRATION NUMBER: 33451

REFERENCE/PACKET NUMBER: Atty File No. 5789542 9301

TELECOMMUNICATION INFORMATION:

TELEPHONE: 504 387-3221

TELEFAX: 504 346-8049

INFORMATION FOR SEQ ID NO: 40:

SEQUENCE CHARACTERISTICS:

LENGTH: 21 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-08-944-133 40

Query Match 75.8%; Score 50; DB 1; Length 21;
 Best Local Similarity 92.3%; Prod. No. 0.28;
 Matches 12; Conservative 0; Mismatches 1; Indels 0;

QY 4 KALKALKALKAL 15
 11111 11111
 DB 5 KALKALKALKAL 17

RESULT 9

US-08-944-133 44

Sequence 44, Application US/08944133

Patent No. 5789542

GENERAL INFORMATION:

APPLICANT: McLaughlin, Mark L

TITLE OF INVENTION: Amphipathic Peptides

NUMBER OF SEQUENCES: 54

CORRESPONDENCE ADDRESS:

ADDRESSEE: John H. Rummels

STREET: P. O. Box 2471

CITY: Baton Rouge

STATE: LA

COUNTRY: USA

ZIP: 70821 2471

COMPUTER PEAKABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: IBM PC compatible

SOFTWARE: Patent in Release #1.0, Version #1.26

us-09-820-053a-43.ra1

Thu Aug 21 08:36:29 2003

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CURRENT APPLICATION DATA: 08/24/94 133
APPLICATION NUMBER: 08/24/94 133
FILING DATE: 06-OCT-1997
CLASSIFICATION: 5530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/789,077
FILING DATE: 03-FEB-1997
APPLICATION NUMBER: US/08/681,075
FILING DATE:
APPLICATION NUMBER: US/98/232,525
FILING DATE: 22-APR-1994
ATTORNEY/AGENT INFORMATION:
NAME: Runnels, John B
REGISTRATION NUMBER: 43451
REFERENCE/SEQUENT NUMBER: 5789542 9301
TELECOMMUNICATION INFORMATION:
TELEPHONE: 504 346-8049
TELEFAX: 504 346-8049
INFORMATION FOR SEQ ID NO: 44:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US 08-944-133-44

Query Match 75.8% Score 50: DB 1: Length 21:
Best local Similarity 92.3% Pred. No. 0.28:
Matches 12: Conservative 0: Mismatches 1: Indels 0: Gaps 0:

QY 3 KALKALKALKAL 15
DB 2 KALKALKALKAL 14

RESULT 10
US-08-241-740A-49
Sequence 49: Application US/08241740A
Patent No. 5561107
GENERAL INFORMATION:
APPLICANT: JAYNES, JESSE M.
APPLICANT: JULIAN, GORDON R.
TITLE OF INVENTION: METHOD OF ENHANCING WOUND HEALING BY STIMULATING FIBROBLAST AN
CORRESPONDENCE ADDRESS:
ADDRESSER: STEVEN J. HULTQUIST
ADDRESS: INTERNATIONAL PROPERTY/TECHNOLOGY LAW
STREET: 200 PARK DRIVE, SUITE 210
CITY: P.O. BOX 14329
STATE: RESEARCH TRIANGLE PARK
COUNTRY: NORTH CAROLINA
ZIP: 27709
COMPUTER READABLE FORM:
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE
COMPUTER: APPLE MACINTOSH
OPERATING SYSTEM: MACINTOSH
SOFTWARE: MS WORD 5.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/06/251,740A
FILING DATE: 04-20-94
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/225,476
FILING DATE: 04-08-94
APPLICATION NUMBER: 08/039,620
FILING DATE: 06-04-93
APPLICATION NUMBER: 08/148,491
FILING DATE: 11-08-93
APPLICATION NUMBER: 08/148,889
FILING DATE: 11-08-93
ATTORNEY/AGENT INFORMATION:
NAME: HULTQUIST, STEVEN J.

CURRENT APPLICATION DATA: 08/24/94 133
APPLICATION NUMBER: 08/24/94 133
FILING DATE: 06-OCT-1997
CLASSIFICATION: 5530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/789,077
FILING DATE: 03-FEB-1997
APPLICATION NUMBER: US/08/681,075
FILING DATE:
APPLICATION NUMBER: US/98/232,525
FILING DATE: 22-APR-1994
ATTORNEY/AGENT INFORMATION:
NAME: Runnels, John B
REGISTRATION NUMBER: 43451
REFERENCE/SEQUENT NUMBER: 5789542 9301
TELECOMMUNICATION INFORMATION:
TELEPHONE: 504 346-8049
TELEFAX: 504 346-8049
INFORMATION FOR SEQ ID NO: 44:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US 08-944-133-44

Query Match 75.8% Score 50: DB 1: Length 21:
Best local Similarity 92.3% Pred. No. 0.28:
Matches 12: Conservative 0: Mismatches 1: Indels 0: Gaps 0:

QY 3 KALKALKALKAL 15
DB 2 KALKALKALKAL 14

RESULT 10
US-08-241-740A-49
Sequence 49: Application US/08241740A
Patent No. 5561107
GENERAL INFORMATION:
APPLICANT: JAYNES, JESSE M.
APPLICANT: JULIAN, GORDON R.
TITLE OF INVENTION: METHOD OF ENHANCING WOUND HEALING BY STIMULATING FIBROBLAST AN
CORRESPONDENCE ADDRESS:
ADDRESSER: STEVEN J. HULTQUIST
ADDRESS: INTERNATIONAL PROPERTY/TECHNOLOGY LAW
STREET: 200 PARK DRIVE, SUITE 210
CITY: P.O. BOX 14329
STATE: RESEARCH TRIANGLE PARK
COUNTRY: NORTH CAROLINA
ZIP: 27709
COMPUTER READABLE FORM:
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE
COMPUTER: APPLE MACINTOSH
OPERATING SYSTEM: MACINTOSH
SOFTWARE: MS WORD 5.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/06/251,740A
FILING DATE: 04-20-94
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/225,476
FILING DATE: 04-08-94
APPLICATION NUMBER: 08/039,620
FILING DATE: 06-04-93
APPLICATION NUMBER: 08/148,491
FILING DATE: 11-08-93
APPLICATION NUMBER: 08/148,889
FILING DATE: 11-08-93
ATTORNEY/AGENT INFORMATION:
NAME: HULTQUIST, STEVEN J.

CURRENT APPLICATION DATA: 28021
REFERENCE/SEQUENT NUMBER: 4312-106
TELECOMMUNICATION INFORMATION:
TELEPHONE: (919)990-9541
TELEFAX: (919)990-9542
INFORMATION FOR SEQ ID NO: 49:
SEQUENCE CHARACTERISTICS:
LENGTH: 23
TYPE: AMINO ACID
TOPOLOGY: LINEAR
MOLECULE TYPE: PEPTIDE
DESCRIPTION: NO
HYPOTHETICAL: NO
FRAGMENT TYPE: COMPLETE PEPTIDE
ORIGINAL SOURCE: SYNTHETIC
IMMEDIATE SOURCE: SYNTHETIC
PUBLICATION INFORMATION: NOT PREVIOUSLY PUBLISHED
US-08-231-730A-49

Query Match 72.7% Score 48: DB 1: Length 23:
Best local Similarity 80.0% Pred. No. 0.58:
Matches 12: Conservative 0: Mismatches 4: Indels 0: Gaps 0:

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKAL 15

RESULT 11
US-08-427-001C-44
Sequence 44: Application US/98427001C
Patent No. 5717064
GENERAL INFORMATION:
APPLICANT: JULIAN, GORDON R.
TITLE OF INVENTION: METHYLATED LYSINE-RICH LYLLI PEPTIDES.
TITLE OF INVENTION: AM. BELL. 0: 552,667 THE SAME IN METHYLATED ALKYLATION
NUMBER OF SEQUENCES: 48
CORRESPONDENCE ADDRESS:
ADDRESSER: ROTHWELL, FLORENCE K. KURZ
STREET: 555 THIRTEENTH STREET, N.W.
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE
COMPUTER: IBM COMPATIBLE
OPERATING SYSTEM: DOS
SOFTWARE: WordPerfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/98/427,001C
FILING DATE: 24-APR-95
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: U.S. 08/148,899
FILING DATE: 08-NOV-93
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: WALKER, BARBARA W.
REGISTRATION NUMBER: 45,400
REFERENCE/SEQUENT NUMBER: 2093 105A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)783-6040
TELEFAX: (202)783-6041
INFORMATION FOR SEQ ID NO: 44:
SEQUENCE CHARACTERISTICS:
LENGTH: 23
TYPE: AMINO ACID
TOPOLOGY: LINEAR
MOLECULE TYPE: PEPTIDE
US-08-427-001C-44

Query Match 72.7% Score 48: DB 1: Length 23:

```


COMPUTER: IBM COMPATIBLE
 OPERATING SYSTEM: DOS
 SOFTWARE: WORDPERFECT 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: 08/279,472
 FILING DATE: 21-JUL-1995
 CLASSIFICATION: 536
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/279,472
 FILING DATE: 22-JUL-1994
 CLASSIFICATION: 536
 ATTORNEY/AGENT INFORMATION:
 NAME: WALKER, BARBARA W.
 REGISTRATION NUMBER: 35,400
 REFERENCE/DOCKET NUMBER: 2093-117A
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (202)783-6040
 TELEFAX: (202)783-6031
 INFORMATION FOR SEQ ID NO: 44:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 23
 TYPE: AMINO ACID
 TOPOLOGY: LINEAR
 MOLECULE TYPE: PEPTIDE
 DESCRIPTION: COMPLETE PEPTIDE
 HYPOTHETICAL: NO
 FRAGMENT TYPE: COMPLETE PEPTIDE
 ORIGINAL SOURCE: SYNTHETIC
 IMMEDIATE SOURCE: SYNTHETIC
 PUBLICATION INFORMATION: NOT PREVIOUSLY PUBLISHED
 US 08-505,486-44

Query Match 72.7% Score 48 DB 2 Length 23;
 Best Local Similarity 80.0% Pred. No. 0.58;
 Matches 12; Conservative 0; Mismatches 4; Indels 0;

QY 1 FAKALKALKAKAL 15
 DB 1 FAKALKALKAKAL 15

RESULT 15
 US-08-689-489C-39
 Sequence 39, Application US/08689489C
 Patent No. 6001805
 GENERAL INFORMATION:
 APPLICANT: JESSE M. JAYNES, Gordon R. Julian
 TITLE OF INVENTION: Method of Enhancing Wound Healing By
 TITLE OF INVENTION: Stimulating Fibroblast and Keratinocyte Growth in
 TITLE OF INVENTION: Vivo, Utilizing Amphipathic Peptides
 NUMBER OF SEQUENCES: 46
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Rothwell, Figg, Ernst & Kurz
 STREET: 555 14TH STREET
 CITY: Washington
 STATE: DC
 COUNTRY: USA
 ZIP: 20004
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC Compatible
 OPERATING SYSTEM: PC DOS/MS-DOS
 SOFTWARE: Patent In Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/689,489C
 FILING DATE: August 12, 1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/231,730
 FILING DATE: April 20, 1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/225,476
 FILING DATE: April 8, 1994
 PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/039,620
 FILING DATE: June 4, 1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/148,889
 FILING DATE: No. 6001805/08/148,889
 PRIOR APPLICATION DATA: 08/148,889
 APPLICATION NUMBER: 08/148,491
 FILING DATE: No. 6001805/08/148,491
 ATTORNEY/AGENT INFORMATION:
 NAME: MARK T. BOWDITCH
 REGISTRATION NUMBER: 40,415
 REFERENCE/DOCKET NUMBER: 2094-120
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-783-6040
 TELEFAX: 202-783-6031
 INFORMATION FOR SEQ ID NO: 39:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 23 amino acids
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: Linear
 MOLECULE TYPE: peptide
 HYPOTHETICAL: NO
 FRAGMENT TYPE: Linear
 US-08-689-489C-39

Query Match 72.7% Score 48 DB 3 Length 24;
 Best Local Similarity 80.0% Pred. No. 0.58;
 Matches 12; Conservative 0; Mismatches 4; Indels 0;

QY 1 FAKALKALKAKAL 15
 DB 1 FAKALKALKAKAL 15

RESULT 16
 US-08-801-028 44
 Sequence 44, Application US/08801028
 Patent No. 6018102
 GENERAL INFORMATION:
 APPLICANT: JUAN GARRARINO
 APPLICANT: JESSE M. JAYNES
 TITLE OF INVENTION: UBIQUITIN-LINKED PEPTIDE FUSION GENES
 NUMBER OF SEQUENCES: 98
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: STEVEN J. BULTOUST
 ADDRESSEE: INTELLECTUAL PROPERTY/TECHNOLOGY LAW
 STREET: 200 PARK DRIVE, SUITE 210
 STREET: P.O. BOX 14329
 CITY: PEASEPORT TRIANGLE PARK
 STATE: NORTH CAROLINA
 COUNTRY: USA
 ZIP: 27709
 COMPUTER READABLE FORM:
 MEDIUM TYPE: DISKETTE, 3.5 INCH 1.4 MB ST-RAP
 COMPUTER: APPLE MACINTOSH
 OPERATING SYSTEM: MACINTOSH
 SOFTWARE: M.S. WORD 5.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: 08/801,028
 FILING DATE: 19 FEB-1997
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/279,472
 FILING DATE: JULY 22, 1994
 APPLICATION NUMBER: 08/225,476
 FILING DATE: 04-20-94
 APPLICATION NUMBER: 08/225,476
 FILING DATE: 04-08-94
 APPLICATION NUMBER: 08/039,620
 FILING DATE: 06-04-94
 APPLICATION NUMBER: 08/148,491
 FILING DATE: 11-08-94


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? FRAGMENT TYPE: LINEAR
? SEQUENCE DESCRIPTION: SEQ ID NO: 39;
US-09-242-802A-49

Query Match
Best Local Similarity 80.0% Score 48; DB 3; Length 23;
Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
Db 1 FALALKALKALKAL 15

RESULT 19
US-09-482-611R-44
? Sequence 44, Application US/09482611R
? Patent No. 6448691
? GENERAL INFORMATION:
? APPLICANT: Belknap, Joan
? TITLE OF INVENTION: Ubiquitin-Lytic Peptide Fusion Gene Constructs, Protein Products
? FILE REFERENCE: 2093-149
? CURRENT APPLICATION NUMBER: US 09-482-611R
? PRIOR FILING DATE: 2000-01-14
? PRIOR APPLICATION NUMBER: US 08/801,028
? PRIOR FILING DATE: 1997-02-19
? PRIOR APPLICATION NUMBER: US 08-2076,479
? NUMBER OF SEQ ID NOS: 102
? SOFTWARE: Patent In version 3.1
? SEQ ID NO 44
? LENGTH: 23
? TYPE: PEPTIDE
? ORGANISM: Artificial Sequence
? OTHER INFORMATION: Lytic Peptide
US-09-482-611R-44

Query Match
Best Local Similarity 80.0% Score 48; DB 4; Length 23;
Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
Db 1 FALALKALKALKAL 15

RESULT 20
US-09-019-922A-44
? Sequence 44, Application US/09019922A
? Patent No. 6559281
? GENERAL INFORMATION:
? APPLICANT: JAYNES, JESSE M.
? TITLE OF INVENTION: NON-NATURALLY OCCURRING SYNTHETIC LYTIC
? FILE REFERENCE: 45
? NUMBER OF SEQUENCES: 45
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: ROTHWELL, FIGG, ERNST & MANECK
? STREET: 555 Thirteenth Street, N.W.
? CITY: Washington
? STATE: D.C.
? COUNTRY: USA
? ZIP: 20004
? COMPUTER READABLE FORM:
? MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE
? COMPUTER: IBM COMPATIBLE
? OPERATING SYSTEM: DOS
? SOFTWARE: WordPad
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US-09-019-922A
? FILING DATE: 06-FEB-98
? CLASSIFICATION:

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? PRIOR APPLICATION DATA:
? APPLICATION NUMBER: U.S.08/427,001
? FILING DATE: 24-APR-95
? PRIOR APPLICATION DATA:
? APPLICATION NUMBER: U.S.08/148,899
? FILING DATE: 08-NOV-94
? ATTORNEY/AGENT INFORMATION:
? NAME: BOWDITCH, MARK L.
? REGISTRATION NUMBER: 40,415
? REFERENCE/EXCERPT NUMBER: 2093-149A
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (202)783-6040
? TELEFAX: (202)783-6041
? INFORMATION FOR SEQ ID NO: 44:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 23
? TYPE: AMINO ACID
? TOPOLOGY: LINEAR
? MOLECULE TYPE: PEPTIDE
US-09-019-922A-44

Query Match
Best Local Similarity 80.0% Score 48; DB 4; Length 23;
Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
Db 1 FALALKALKALKAL 15

RESULT 21
PCT US94-12550-47
? Sequence 47, Application PCT/US9412550
? GENERAL INFORMATION:
? APPLICANT: JULIAN, GORDON R.
? TITLE OF INVENTION: METHYLATED LYSINE RICH LYTIC
? FILE REFERENCE: PEPTIDES AND METHOD OF
? TITLE OF INVENTION: MAKING SAME BY REDUCTIVE
? NUMBER OF SEQUENCES: 47
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: STEVEN J. HOLLOUIS
? STREET: 200 PARK DRIVE, SUITE 210
? CITY: RESEARCH TRIANGLE PARK
? STATE: NORTH CAROLINA
? COUNTRY: USA
? ZIP: 27709
? COMPUTER READABLE FORM:
? MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE
? COMPUTER: APPLE MACINTOSH
? OPERATING SYSTEM: MACINTOSH
? SOFTWARE: M.S. WORD 5.0
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: PCT/US94/12550
? FILING DATE: NOVEMBER 8, 1994
? PRIOR APPLICATION DATA: NONE
? ATTORNEY/AGENT INFORMATION:
? NAME: HOLLOUIS, STEVEN J.
? REGISTRATION NUMBER: 28021
? REFERENCE/EXCERPT NUMBER: 4013-101
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (919)990-9541
? TELEFAX: (919)990-9542
? INFORMATION FOR SEQ ID NO: 47:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 23
? TYPE: AMINO ACID
? TOPOLOGY: LINEAR
? MOLECULE TYPE: PEPTIDE
? DESCRIPTION: PEPTIDE
? HYPOTHETICAL: NO

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3 SOFTWARE: DW4 V2
3 CURRENT APPLICATION DATA:
3 APPLICATION NUMBER: US/07/909,455A
3 FILING DATE: 19920702
3 CLASSIFICATION: 514
3 PRIOR APPLICATION DATA:
3 APPLICATION NUMBER: US 07686115
3 FILING DATE: 15 APR 1991
3 APPLICATION NUMBER: US 07476629
3 FILING DATE: 08 FEB 1990
3 ATTORNEY/AGENT INFORMATION:
3 NAME: Elstein, Elliot M.
3 REGISTRATION NUMBER: 24,025
3 REFERENCE/BOOKLET NUMBER: 421,250-122
3 TELECOMMUNICATION INFORMATION:
3 TELEPHONE: 201-994-1744
3 TELEFAX: 201-994-1700
3 INFORMATION FOR SEQ ID NO: 8:
3 SEQUENCE CHARACTERISTICS:
3 LENGTH: 21 amino acids
3 TYPE: AMINO ACID
3 STRANDEDNESS:
3 TOPOLOGY: linear
3 MOLECULE TYPE: peptide
3 FEATURE:
3 OTHER INFORMATION: amide terminated
3 US 07-908 455A R

3 Query Match: 60.6%, Score 40, DB 1, Length 21;
3 Best Local Similarity: 61.5%, Prod. No. 7, 4;
3 Matches: 8; Conservative: 4; Mismatches: 1; Indels: 0; Gaps: 0;

3 QY 3 KAIKAIKAIKAI 15
3 DB 5 KAIKAIKAIKAI 17

3 RESULT 28
3 US 08-444 120-44
3 Sequence 34, Application US/08/444,120
3 Patent No. 5466479
3 GENERAL INFORMATION:
3 APPLICANT: Baker, Margaret A.
3 APPLICANT: Jacob, Leonard S.
3 APPLICANT: Maloy, W. Lee
3 TITLE OF INVENTION: Treatment of Gynecological
3 TITLE OF INVENTION: Malignancies with
3 TITLE OF INVENTION: Biologically Active Peptides
3 NUMBER OF SEQUENCES: 117
3 CORRESPONDENCE ADDRESS:
3 ADDRESSEE: Carolina, Pyrie, Bitt, Gilliland,
3 STREET: 6 Becker Farm Road
3 CITY: Roseland
3 STATE: New Jersey
3 COUNTRY: USA
3 ZIP: 07068
3 COMPUTER READABLE FORM:
3 MEDIUM TYPE: 3.5 inch diskette
3 COMPUTER: IBM PS/2
3 OPERATING SYSTEM: PC-DOS
3 SOFTWARE: DW4 V2
3 CURRENT APPLICATION DATA:
3 APPLICATION NUMBER: US/08/444,120
3 FILING DATE:
3 CLASSIFICATION: 514
3 PRIOR APPLICATION DATA:
3 APPLICATION NUMBER: US/08/297,950
3 FILING DATE:
3 APPLICATION NUMBER: US/08/226,108
3 FILING DATE:
3 APPLICATION NUMBER: US/07/987,462
3 FILING DATE:

```

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3 ATTORNEY/AGENT INFORMATION:
3 NAME: Elstein, Elliot M.
3 REGISTRATION NUMBER: 24,025
3 REFERENCE/BOOKLET NUMBER: 421,250-122
3 TELECOMMUNICATION INFORMATION:
3 TELEPHONE: 201-994-1700
3 TELEFAX: 201-994-1744
3 INFORMATION FOR SEQ ID NO: 4:
3 SEQUENCE CHARACTERISTICS:
3 LENGTH: 21 amino acids
3 TYPE: amino acid
3 STRANDEDNESS:
3 TOPOLOGY: linear
3 MOLECULE TYPE: peptide
3 US 08-444 120-44

3 Query Match: 60.6%, Score 40, DB 1, Length 21;
3 Best Local Similarity: 61.5%, Prod. No. 7, 4;
3 Matches: 8; Conservative: 4; Mismatches: 1; Indels: 0; Gaps: 0;

3 QY 3 KAIKAIKAIKAI 15
3 DB 5 KAIKAIKAIKAI 17

3 RESULT 29
3 US 08-465 425-44
3 Sequence 34, Application US/08/465,425
3 Patent No. 6066564
3 GENERAL INFORMATION:
3 APPLICANT: Maudlin Pharmaceuticals Inc.
3 APPLICANT: 5110 Campus Drive
3 APPLICANT: Edgewood, Georgia, GA 30022
3 TITLE OF INVENTION: Biologically Active Peptides
3 TITLE OF INVENTION: N-Terminal Substitutions
3 NUMBER OF SEQUENCES: 154
3 CORRESPONDENCE ADDRESS:
3 ADDRESSEE: Fluorophore Bioscience Research &
3 ADDRESSEE: Inc.
3 STREET: 4000 E. Street, N.W. Suite 200
3 CITY: Washington
3 STATE: DC
3 COUNTRY: USA
3 ZIP: 20006-3915
3 COMPUTER READABLE FORM:
3 MEDIUM TYPE: Floppy Disk
3 COMPUTER: IBM PC compatible
3 OPERATING SYSTEM: PC-DOS/MS-DOS
3 SOFTWARE: Patent In Release #120, Version #1.0
3 CURRENT APPLICATION DATA:
3 APPLICATION NUMBER: US/08/465,425
3 FILING DATE: 05 JUN 1999
3 CLASSIFICATION: 514
3 PRIOR APPLICATION DATA:
3 APPLICATION NUMBER: 08/184,462
3 FILING DATE: 18 JAN 94
3 PRIOR APPLICATION DATA:
3 APPLICATION NUMBER: 07/994,200
3 FILING DATE: 01 JUN 92
3 ATTORNEY/AGENT INFORMATION:
3 NAME: Fortis, John W.
3 REGISTRATION NUMBER: 51,064
3 REFERENCE/BOOKLET NUMBER: 092,000
3 TELECOMMUNICATION INFORMATION:
3 TELEPHONE: (202) 408-4400
3 TELEFAX: (202) 408-4400
3 INFORMATION FOR SEQ ID NO: 4:
3 SEQUENCE CHARACTERISTICS:
3 LENGTH: 21 amino acids
3 TYPE: amino acid
3 TOPOLOGY: linear
3 MOLECULE TYPE: peptide
3 US 08-465 425-44

```



```

1 PRIOR APPLICATION DATA:
2 APPLICATION NUMBER: 08/225,476
3 FILING DATE: 04-08-94
4 APPLICATION NUMBER: 08/049,420
5 FILING DATE: 06-04-94
6 APPLICATION NUMBER: 08/148,491
7 FILING DATE: 11-08-94
8 APPLICATION NUMBER: 08/148,889
9 FILING DATE: 11-08-94
10 ATTORNEY/AGENT INFORMATION:
11 NAME: HILTON/ST. STEVEN J.
12 REGISTRATION NUMBER: 28021
13 REFERENCE/PACKET NUMBER: 4015 106
14 TELECOMMUNICATION INFORMATION:
15 TELEPHONE: (919)990-9541
16 TELEFAX: (919)990-9542
17 INFORMATION FOR SEQ ID NO: 42:
18 SEQUENCE CHARACTERISTICS:
19 LENGTH: 24
20 TYPE: AMINO ACID
21 TOPOLOGY: LINEAR
22 MOLECULE TYPE:
23 DESCRIPTION: PEPTIDE
24 HYPOTHETICAL: NO
25 FRAGMENT TYPE: COMPLETE PEPTIDE
26 ORIGINAL SOURCE: SYNTHETIC
27 IMMEDIATE SOURCE: SYNTHETIC
28 PUBLICATION INFORMATION: NOT PREVIOUSLY PUBLISHED
29 US 08 231-740A-42

```

Query Match 59.1% Score 492 18 17 Length 24

Best Local Similarity 53.9% Prod. No. 112

MATCHES 92 Conservative 4 Mismatches 3 Indels 0 Caps 0

QV 1 FAKAKALKAKKAL 15

11 1111 1111

1b 1 FAKAKALKAKKAK 15

RESULT 34

US 08 505 486 47

Sequence 47, Application US/0805486

Patent No. 5455574

GENERAL INFORMATION:

APPLICANT: JESSE M. JAYNES

TITLE OF INVENTION: ORBICULIN-LYTIC PEPTIDE FUSION GENE

TITLE OF INVENTION: CONSTRUCTS, PROTEIN PRODUCTS, PEPTIDE

TITLE OF INVENTION: METHODS OF MAKING AND USING SAME

NUMBER OF SEQUENCES: 98

CORRESPONDENCE ADDRESS:

ADDRESS: ROTHWELL, FLOD, ERNST & KURZ

STREET: 555 THIRTEENTH STREET N.W.

CITY: WASHINGTON

STATE: D. C.

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB STORAGE

COMPUTER: IBM COMPATIBLE

OPERATING SYSTEM: DOS

SOFTWARE: WORDPERFECT 5.1

CURRENT APPLICATION DATA:

APPLICATION NUMBER: 08/095,486

FILING DATE: 21-JUL-1995

CLASSIFICATION: 536

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/092,754

FILING DATE: 22-JUL-1994

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: WALKER, BARBARA W.

REGISTRATION NUMBER: 45,400

REFERENCE/PACKET NUMBER: 2099 117A

TELECOMMUNICATION INFORMATION:

TELEPHONE: (202)783 6040

TELEFAX: (202)783 6041

INFORMATION FOR SEQ ID NO: 43:

SEQUENCE CHARACTERISTICS:

LENGTH: 24

TYPE: AMINO ACID

TOPOLOGY: LINEAR

MOLECULE TYPE:

DESCRIPTION: PEPTIDE

HYPOTHETICAL: NO

FRAGMENT TYPE: COMPLETE PEPTIDE

ORIGINAL SOURCE: SYNTHETIC

IMMEDIATE SOURCE: SYNTHETIC

PUBLICATION INFORMATION: NOT PREVIOUSLY PUBLISHED

US-08-505 486 47

Query Match 59.1%

Best Local Similarity 53.9% Prod. No. 112

MATCHES 92 Conservative 4 Mismatches 3 Indels 0 Caps 0

QV 1 FAKAKALKAKKAL 15

11 1111 1111

1b 1 FAKAKALKAKKAK 15

RESULT 34

US 08 689 486 47

Sequence 47, Application US/0806486

Patent No. 6001806

GENERAL INFORMATION:

APPLICANT: JESSE M. JAYNES, GEORGE F. JAYNES

TITLE OF INVENTION: ORBICULIN-LYTIC PEPTIDE FUSION GENE

TITLE OF INVENTION: CONSTRUCTS, PROTEIN PRODUCTS, PEPTIDE

TITLE OF INVENTION: METHODS OF MAKING AND USING SAME

NUMBER OF SEQUENCES: 46

CORRESPONDENCE ADDRESS:

ADDRESS: ROTHWELL, FLOD, ERNST & KURZ

STREET: 555 14TH STREET

CITY: WASHINGTON

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: FLOPPY DISK

COMPUTER: IBM PC COMPATIBLE

OPERATING SYSTEM: DOS/MS-DOS

SOFTWARE: PATENT IN REPOSE #1.0, Version #1.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/069,486

FILING DATE: August 12, 1990

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/069,486

FILING DATE: April 20, 1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/225,476

FILING DATE: April 8, 1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/069,486

FILING DATE: June 4, 1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/148,889

FILING DATE: Not Known/Unknown, 1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/148,491

FILING DATE: Not Known/Unknown, 1994

NAME: WALKER, BARBARA W.

REGISTRATION NUMBER: 45,400

TELECOMMUNICATION INFORMATION:

TELEPHONE: 202 783 6040

QY 1 FAKALKALKALKAL 15
 1111111111
 DB 1 FAIAIKAIKAIKAI 15

RESULT 47

US 09 242 802A 42

Sequence 47, Application US/2002/02802A

Patent No. 6191110

GENERAL INFORMATION:

APPLICANT: Jesse M. Jaynes, Gordon R. Julian

TITLE OF INVENTION: Method of Enhancing Wound Healing by
 Stimulating Fibroblast and Keratinocyte Growth in
 Vivo, Utilizing Amphipathic Peptides

NUMBER OF SEQUENCES: 46

CORRESPONDENCE ADDRESS:

ADDRESSEE: Redwood, Figg, Ernst & Marbeck

STREET: 555 14TH STREET

CITY: Washington

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC Compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: 08/689,489

FILING DATE: 19-Jan-1999

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/689,489

FILING DATE: August 12, 1996

APPLICATION NUMBER: 08/223,730

FILING DATE: April 20, 1994

APPLICATION NUMBER: 08/225,476

FILING DATE: April 8, 1994

APPLICATION NUMBER: 08/049,620

FILING DATE: June 4, 1993

APPLICATION NUMBER: 08/148,889

FILING DATE: No. 6191110, October 8, 1993

APPLICATION NUMBER: 08/148,491

FILING DATE: No. 6191110, October 9, 1993

ATTORNEY/AGENT INFORMATION:

NAME: Mark L. Bowditch

REGISTRATION NUMBER: 40,415

REFERENCE/WORK NUMBER: 2093 142

TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-783-6040

TELEFAX: 202-783-6041

INFORMATION FOR SEQ ID NO: 42:

SEQUENCE CHARACTERISTICS:

LENGTH: 23 amino acids

TYPE: amino acid

STRANDNESS: unknown

TOPOLOGY: linear

MOLECULE TYPE: peptide

HYPOTHETICAL: NO

FRAGMENT TYPE: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 42:

US-09 242 802A 42

Query Match 59.18; Score 49; DB 4; Length 23;
 Best Local Similarity 53.9; Pred. No. 11;
 Matches 8; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15

1111111111

DB 1 FAIAIKAIKAIKAI 15

RESULT 48

US 09 482-611B 47

Sequence 47, Application US/2004/06111B

Patent No. 6448491

GENERAL INFORMATION:

APPLICANT: Garbath, William

TITLE OF INVENTION: Dopamine Receptor Antagonists

TITLE OF INVENTION: The Form, and Methods of Making and Using

FILE REFERENCE: 2004 149

CURRENT APPLICATION NUMBER: US/09/482,611B

CURRENT FILING DATE: 2004 01 14

PRIOR APPLICATION NUMBER: US 09/001,028

PRIOR FILING DATE: 1997 02 19

PRIOR ATTORNEY NUMBER: US 09/279,472

PRIOR FILING DATE: 1994 07 22

NUMBER OF SEQ ID NOS: 102

SOFTWARE: Patent in version 4.3.1

SEQ ID NO 47

LENGTH: 23

TYPE: PEI

ORGANISM: Artificial Sequence

FEATURE:

OTHER INFORMATION: Tyrosine Peptide

US 09 482-611B 47

Query Match 59.18; Score 49; DB 4; Length 23

Best Local Similarity 53.9; Pred. No. 11;

Matches 8; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15

1111111111

DB 1 FAIAIKAIKAIKAI 15

RESULT 49

PT 0505 0471B 42

Sequence 42, Application PT/2005/0471B

GENERAL INFORMATION:

APPLICANT: LEMTER BUCHENHOEFER, LTD.

TITLE OF INVENTION: METHOD OF ENHANCING WOUND HEALING BY COMBINING A PEPTIDE

NUMBER OF SEQUENCES: 46

CORRESPONDENCE ADDRESS:

ADDRESSEE: FRANK WASSEMAN

ADDRESS: INDUSTRIAL TRACTIVITY/TECHNICAL LAW

STREET: 200 LARK DRIVE, SUITE 210

CITY: P.O. BOX 14429

STATE: NORTH CAROLINA

COUNTRY: USA

ZIP: 27709

COMPUTER READABLE FORM:

MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.4 MB 51+KB

COMPUTER: APPLE MACINTOSH

OPERATING SYSTEM: MACINTOSH

SOFTWARE: M.S. WORD 6.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: PT/2005/0471B

FILING DATE:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/271,750

FILING DATE: 20 04 94

ATTORNEY/AGENT INFORMATION:

NAME: WASSEMAN, FRANK S.

REGISTRATION NUMBER: 44273

REFERENCE/WORK NUMBER: 4013 107

TELECOMMUNICATION INFORMATION:

TELEPHONE: (919) 990 9541

TELEFAX: (919) 990 9542

INFORMATION FOR SEQ ID NO: 42:

SEQUENCE CHARACTERISTICS:

LENGTH: 23

TYPE: AMINO-ACID

TOPOLOGY: LINEAR


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1 STRANDEDNESS: Single
2 Topology: Linear
3 US 08 944 434-43
4
5 Query Match
6 Best local similarity 57.68; Score 48; DB 1; Length 14;
7 Matches 0; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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10 4 KALKALKALK 12
11 11111111
12 4 KALKALKALK 13
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14
15 RESULT 44
16 US 08 944 434-29
17 Sequence 29, Application US/08944133
18 Patent No. 5789942
19 GENERAL INFORMATION:
20 APPLICANT: McLaughlin, Mark L
21 APPLICANT: Becker, Calvin L
22 TITLE OF INVENTION: Amphipathic Peptides
23 NUMBER OF SEQUENCES: 54
24 CORRESPONDENCE ADDRESS:
25 ADDRESSEE: John H. Rannels
26 STREET: P. O. Box 2471
27 CITY: Baton Rouge
28 STATE: LA
29 COUNTRY: USA
30 ZIP: 70821 2471
31
32 COMPUTER READABLE FORM:
33 MEDIUM TYPE: Floppy disk
34 COMPUTER: IBM PC compatible
35 OPERATING SYSTEM: PC DOS/MS DOS
36 SOFTWARE: Patented Release #1.0, Version #1.25
37
38 CURRENT APPLICATION DATA:
39 APPLICATION NUMBER: US/98/044,133
40 FILING DATE: 06-OCT-1997
41 CLASSIFICATION: 5540
42 PRIOR APPLICATION DATA:
43 APPLICATION NUMBER: 08/789,077
44 FILING DATE: 03-FEB-1997
45 APPLICATION NUMBER: 08/200,403
46 FILING DATE:
47 APPLICATION NUMBER: 08/232,525
48 FILING DATE: 22-APR-1994
49 ATTORNEY/AGENT INFORMATION:
50 NAME: Rannels, John H
51 REGISTRATION NUMBER: 3451
52 REFERENCE/CKET NUMBER: Atty File No. 5789942 9401
53 TELECOMMUNICATION INFORMATION:
54 TELEPHONE: 504 487-4221
55 TELEFAX: 504 446 8049
56 INFORMATION FOR SEQ ID NO. 29:
57 LENGTH: 14 amino acids
58 TYPE: amino acid
59 STRANDEDNESS: single
60 TOPOLOGY: Linear
61 MOLECULE TYPE: peptide
62
63 US 08 944 434-29
64
65 Query Match
66 Best local similarity 61.88; Score 48; DB 1; Length 14;
67 Matches 0; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
68
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70 4 KALKALKALK 13
71 11111111
72 1 KALKALKALK 11
73
74
75 RESULT 44
76 US 07 908 455A 7
77 Sequence 7, Application US/07908455A

```

```

1 Patent No. 5450247
2 GENERAL INFORMATION:
3 APPLICANT: Berkowitz, Barry A.
4 APPLICANT: Kati, D. Prasad
5 APPLICANT: Mahoy, W. Lee
6 TITLE OF INVENTION: Novel Cysteine-Modified Peptides
7 FIELD OF INVENTION: Basic Therapeutics
8 NUMBER OF SEQUENCES: 69
9 CORRESPONDENCE ADDRESS:
10 ADDRESSEE: Catechi, Rymos, Baton Rouge, Louisiana
11 ADDRESSEE: Catechi & Stewart
12 STREET: 6 Becker Farm Road
13 CITY: Roseland
14 STATE: New Jersey
15 COUNTRY: USA
16 ZIP: 07068
17
18 COMPUTER READABLE FORM:
19 MEDIUM TYPE: 3.5 inch diskette
20 COMPUTER: IBM PS/2
21 OPERATING SYSTEM: PC DOS
22 SOFTWARE: IBM V2
23
24 CURRENT APPLICATION DATA:
25 APPLICATION NUMBER: US/97/908,455A
26 FILING DATE: 19920762
27 CLASSIFICATION: 514
28 PRIOR APPLICATION DATA:
29 APPLICATION NUMBER: US 07606115
30 FILING DATE: 15-APR-1991
31 APPLICATION NUMBER: US 07466099
32 FILING DATE: 08-FEB-1990
33 ATTORNEY/AGENT INFORMATION:
34 NAME: Catechi, Elliot M.
35 REGISTRATION NUMBER: 34,525
36 REFERENCE/CKET NUMBER: 421,001,122
37 TELECOMMUNICATION INFORMATION:
38 TELEFAX: 201 994 1744
39 TELEPHONE: 201 994 1700
40 INFORMATION FOR SEQ ID NO. 7:
41 SEQUENCE CHARACTERISTICS:
42 LENGTH: 21 amino acids
43 TYPE: AMINO ACID
44 COMPLETENESS:
45 TOPOLOGY: Linear
46 MOLECULE TYPE: peptide
47 FEATURE:
48 OTHER INFORMATION: amino-terminated
49
50 US 07 908 455A 7
51
52 Query Match
53 Best local similarity 62.99; Score 48; DB 1; Length 21;
54 Matches 10; Conservative 0; Mismatches 11; Indels 0; Gaps 0;
55
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57 4 KALKALKALKAL 15
58 1111111111
59 1 KALKALKALKAL 14
60
61
62 RESULT 45
63 US 08 434 120-43
64 Sequence 43, Application US/08434120
65 Patent No. 563449
66 GENERAL INFORMATION:
67 APPLICANT: Becker, Margaret A.
68 APPLICANT: Jacob, Donald S.
69 APPLICANT: Mahoy, W. Lee
70 TITLE OF INVENTION: Treatment of Rheumatoid Arthritis
71 FIELD OF INVENTION: Medication with
72 FIELD OF INVENTION: Biotechnology
73 NUMBER OF SEQUENCES: 17
74 CORRESPONDENCE ADDRESS:
75 ADDRESSEE: Catechi, Rymos, Baton Rouge, Louisiana
76 ADDRESSEE: Catechi & Stewart
77 STREET: 6 Becker Farm Road

```

```

1 CITY: Roseland
2 STATE: New Jersey
3 COUNTRY: USA
4 ZIP: 07068
5 COMPUTER READABLE FORM:
6 MEDIUM TYPE: 3.5 inch diskette
7 COMPUTER: IBM PS/2
8 OPERATING SYSTEM: PC-DOS
9 SOFTWARE: DW4.V2
10 CURRENT APPLICATION DATA:
11 APPLICATION NUMBER: US/08/434,120
12 FILING DATE:
13 CLASSIFICATION: 514
14 PRIOR APPLICATION DATA:
15 APPLICATION NUMBER: US/08/297,950
16 FILING DATE:
17 APPLICATION NUMBER: US/08/226,108
18 FILING DATE:
19 APPLICATION NUMBER: US/07/937,462
20 FILING DATE:
21 ATTORNEY/AGENT INFORMATION:
22 NAME: Gustaf, Elliot M.
23 REGISTRATION NUMBER: 24,025
24 REFERENCE/DOCKET NUMBER: 421250-194
25 TELECOMMUNICATION INFORMATION:
26 TELEPHONE: 201-994-1700
27 TELEFAX: 201-994-1744
28 INFORMATION FOR SEQ ID NO: 33:
29 SEQUENCE CHARACTERISTICS:
30 LENGTH: 21 amino acids
31 TYPE: amino acid
32 STRANDNESS:
33 TOPOLOGY: linear
34 MOLECULE TYPE: peptide
35 US-08-434-120-33

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Query Match      57.6%; Score 38; DB 1; Length 21;
Best Local Similarity 76.9%; Pred. No. 14;
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY      4 KALKALKALKAL 15
        | | | | | | | |
Db      1 KIKKALKKLLKAL 13

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Search completed: August 21, 2003, 09:22:34
Job time : 29 secs

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Genware version 5.1.1.6
Copyright (c) 1995-2003 Compugen Ltd

om protein protein search, using sw model

Run on: August 21, 2003, 08:06:46 : Search time 82 seconds
(without alignments)
29.035 Million cell updates/sec

Title: US-09-820-053a-43
Perfect score: 66
Sequence: 1 PARALELLELIZAL 15

Scoring table: BLASTSUM62
Gap 10.0 : Gapext 0.5

Searched: 1107863 seqs, 15822572 residues
total number of hits satisfying chosen parameters: 434095

Minimum DB seq length: 0
Maximum DB seq length: 25

Post processing: Minimum Match ok
Maximum Match 100%
Listed first 45 summaries

Database :	Accession	Length	* Score	Match	length	DB	10	Description
1	AA1985.001	15	24	AA1985.001				Reactive synthet
2	AA1985.001	15	24	AA1985.001				Reactive synthet
3	AA1985.001	15	24	AA1985.001				Cancer treatment, a
4	AA1985.001	15	24	AA1985.001				Basic (positively)
5	AA1985.001	15	24	AA1985.001				Amphiphilic ion ch
6	AA1985.001	15	24	AA1985.001				C-terminal substid
7	AA1985.001	15	24	AA1985.001				Amphiphilic peptid
8	AA1985.001	15	24	AA1985.001				Biologically active
9	AA1985.001	15	24	AA1985.001				Ion channel termin

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	length	DB	10	Description
1	66	100.0	15	24	AA1985.001	Reactive synthet
2	63	90.9	15	24	AA1985.001	Reactive synthet
3	61	77.3	21	15	AA1985.001	Cancer treatment, a
4	61	77.3	21	14	AA1985.001	Basic (positively)
5	61	77.3	21	14	AA1985.001	Amphiphilic ion ch
6	61	77.3	21	14	AA1985.001	C-terminal substid
7	61	77.3	21	14	AA1985.001	Amphiphilic peptid
8	61	77.3	21	14	AA1985.001	Biologically active
9	61	77.3	21	15	AA1985.001	Ion channel termin

10	51	77.3	21	15	AA1985.001	AA1985.001
11	51	77.3	21	15	AA1985.001	AA1985.001
12	51	77.3	21	15	AA1985.001	AA1985.001
13	51	77.3	21	15	AA1985.001	AA1985.001
14	51	77.3	21	15	AA1985.001	AA1985.001
15	51	77.3	21	20	AA1985.001	AA1985.001
16	51	77.3	21	23	AA1985.001	AA1985.001
17	51	77.3	21	19	AA1985.001	AA1985.001
18	51	77.3	21	19	AA1985.001	AA1985.001
19	51	77.3	21	19	AA1985.001	AA1985.001
20	51	77.3	21	19	AA1985.001	AA1985.001
21	51	77.3	21	19	AA1985.001	AA1985.001
22	48	72.7	23	11	AA1985.001	AA1985.001
23	48	72.7	23	14	AA1985.001	AA1985.001
24	48	72.7	23	16	AA1985.001	AA1985.001
25	48	72.7	23	17	AA1985.001	AA1985.001
26	48	72.7	23	17	AA1985.001	AA1985.001
27	48	72.7	23	19	AA1985.001	AA1985.001
28	48	72.7	23	19	AA1985.001	AA1985.001
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58	48	72.7	23	24	AA1985.001	AA1985.001
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65	48	72.7	23	24	AA1985.001	AA1985.001
66	48	72.7	23	24	AA1985.001	AA1985.001
67	48	72.7	23	24	AA1985.001	AA1985.001
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69	48	72.7	23	24	AA1985.001	AA1985.001
70	48	72.7	23	24	AA1985.001	AA1985.001
71	48	72.7	23	24	AA1985.001	AA1985.001
72	48	72.7	23	24	AA1985.001	AA1985.001
73	48	72.7	23	24	AA1985.001	AA1985.001
74	48	72.7	23	24	AA1985.001	AA1985.001
75	48	72.7	23	24	AA1985.001	AA1985.001
76	48	72.7	23	24	AA1985.001	AA1985.001
77	48	72.7	23	24	AA1985.001	AA1985.001
78	48	72.7	23	24	AA1985.001	AA1985.001
79	48	72.7	23	24	AA1985.001	AA1985.001
80	48	72.7	23	24	AA1985.001	AA1985.001
81	48	72.7	23	24	AA1985.001	AA1985.001
82	48	72.7	23	24	AA1985.001	AA1985.001
83	48	72.7	23	24	AA1985.001	AA1985.001
84	48	72.7	23	24	AA1985.001	AA1985.001
85	48	72.7	23	24	AA1985.001	AA1985.001
86	48	72.7	23	24	AA1985.001	AA1985.001
87	48	72.7	23	24	AA1985.001	AA1985.001
88	48	72.7	23	24	AA1985.001	AA1985.001
89	48	72.7	23	24	AA1985.001	AA1985.001
90	48	72.7	23	24	AA1985.001	AA1985.001
91	48	72.7	23	24	AA1985.001	AA1985.001
92	48	72.7	23	24	AA1985.001	AA1985.001
93	48	72.7	23	24	AA1985.001	AA1985.001
94	48	72.7	23	24	AA1985.001	AA1985.001
95	48	72.7	23	24	AA1985.001	AA1985.001
96	48	72.7	23	24	AA1985.001	AA1985.001
97	48	72.7	23	24	AA1985.001	AA1985.001
98	48	72.7	23	24	AA1985.001	AA1985.001
99	48	72.7	23	24	AA1985.001	AA1985.001
100	48	72.7	23	24	AA1985.001	AA1985.001

XX WPI: 2003 221247/21.
 XX
 XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
 PT aene, inhibiting growth of microbial cells, or promoting proliferation
 PI of cells, comprises phenylalanine, leucine, alanine or lysine residues
 XX
 XX Claim 7: Page 6; 13pp; English.
 XX
 XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues, the peptides of the
 CC invention have antibacterial, tumoricide, cytostatic, and vulnerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC aene, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in APP00759, ABR00923 represent the bisactive
 CC peptides of the invention.
 XX
 XX Sequence: 15 AA;
 SQ

Query Match 100.0%; Score 66; DB 24; Length 15;
 Best Local Similarity 100.0%; Pred. No. 0.00094;
 Matches 15; Conservation 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
 DB 1 FAKALKALKALKAL 15

RESULT 2
 ABR00910
 ID ABR00910 standard; peptide: 15 AA.

XX ABR00910;
 AC
 DT 03 APR 2003 (first entry)
 XX
 DE Bioactive synthetic peptide FIAK17CV.

XX
 KW Antibacterial; tumoricide; cytostatic; vulnerary; cancer; cystic fibrosis;
 KW aene; antimicrobial; human fibroblast; human lymphocyte; wound healing;
 KW bioactive.
 XX
 OS Synthetic.

XX Key Location/Qualifiers
 PU Modified-site 15 "c-terminal amide"
 FT W12027-430 A2;
 XX
 XX 19 OCT 2002;
 XX

XX 28 MAR 2002; 2002W0509534;
 XX
 XX 28 MAR 2001; 2001US-279505P;
 PR 28 MAR 2001; 2001US-0820053;
 XX
 XX (HELIX) HELIX BIOMEDIX INC.

XX OWN DR;

XX WPI: 2003 221247/21.

XX
 XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
 PT aene, inhibiting growth of microbial cells, or promoting proliferation
 PI of cells, comprises phenylalanine, leucine, alanine or lysine residues
 XX
 XX Claim 7: Page 9; 13pp; English.
 PS

XX
 CC the invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues, the peptides of the
 CC invention have antibacterial, tumoricide, cytostatic, and vulnerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC aene, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in Appendix ABR00910 represent the bisactive
 CC peptides of the invention.
 XX
 XX Sequence: 15 AA;
 SQ

Query Match 100.0%; Score 66; DB 24; Length 15;
 Best Local Similarity 100.0%; Pred. No. 0.00094;
 Matches 15; Conservation 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AKALKALKALKAL 15
 DB 2 AKALKALKALKAL 15

RESULT 3
 AAR68967
 ID AAR68967 standard; peptide: 20 AA.

XX AAR68967;
 AC
 DT 25 MAR 2003 (updated)
 DT 20 APR 1995 (first entry)
 XX
 DE Cancer treating, amphiphilic ion channel forming peptide; cancer treatment;
 XX
 KW Amphiphilic ion channel forming peptide; cancer treatment;
 KW protease inhibitors.

XX Synthetic;
 XX W09419469-A1;
 PR 01 SEP 1994;
 XX
 XX 22 FEB 1994; 94WO 0302723;
 XX
 XX 26 FEB 1993; 94US 0021602;
 XX
 XX (MAGNA) MAGNAN PHARM INC.

XX Berlyn M, Jacob LS, Mulloy WL;
 XX WPI: 1994 294269/06;
 XX
 XX Treating cancerous growths by administering a peptide and a
 XX peptide(s) and protease inhibitors.
 XX
 XX Claim 2: Page 67; 124pp; English.

XX
 XX AAR68967 to AAR68967 and AAR68967 to AAR68967 are the same peptide
 CC ion channel forming peptides conforming to the same amino acid
 CC sequence, used in combination with one or more additional
 CC inhibitors and other amphiphilic ion channel forming peptide
 CC or proteins; they are effective in the treatment of cancerous
 CC growths, in particular during surgery and following surgery,
 CC they may be useful in inhibiting proteolysis and in promoting
 CC potential "loose" mutant cells capable of forming a tumor
 CC sites.
 CC (Updated on 25 MAR 2003 to correct IN field).

XX Sequence: 20 AA;

Query Match 100.0%; Score 66; DB 24; Length 20;
 Best Local Similarity 85.0%; Pred. No. 0.034;

Matches 12: Conservation 1: Mismatches 1: Indels 0: Gaps 0:

27 2 AKALAKALKAL 15
 10 AAK45054 standard: peptide: 21 AA.
 XX
 AC AAK45054;
 XX
 DT 25-MAR-2003 (updated)
 DT 26 JUN 1994 (first entry)
 XX
 XX Basic (positively charged) polypeptide for N terminal
 DE lipophilic substitution.
 KW ion channel; madarin; RNA; XRP; GTP; coenzyme; sarcosine;
 KW amphiphilic; hydrophobic; hydrophilic; lipophilic; growth;
 KW inhibition; target cell; virus; virally-infected cell;
 KW antitumour; antiviral; antitumor; antiparasitic;
 KW spermicide; wound healing; burn; infection.
 XX
 OS Synthetic.
 XX
 ON W09424138-A1.
 XX
 PD 09-DEC-1993.
 XX
 PF 27 MAY 1993; 94W0 US05192.
 XX
 PR 01 JUN 1993; 92W0 US91201.
 XX
 PA (MAGA) MAGAININ PHARM INC.
 XX
 PI Karl B;
 XX
 PR 1993-405419/50.
 XX
 XX Peptide(s) or proteins with an N-terminal lipophilic substit. -
 PI used for inhibiting growth of target cell, virus or
 PI virally-infected cell
 XX
 PS Disclosure: Page 75-88; 113pp; English.
 XX
 CC A novel compsn. for inhibiting growth of a target cell, virus or
 CC virally-infected cell comprises a peptide of formula T-N(W)-X (1),
 CC X is a biologically active amphiphilic ion channel-forming peptide
 CC or protein; pref. a magainin peptide, a pGLa peptide, a XRP
 CC peptide, a CPP peptide, a coetropin or a sarcotoxin.
 CC N is the nitrogen of the N-terminal amino group.
 CC T is a lipophilic moiety; pref. R-50, where R is a 2-hex
 CC hydrocarbon (alkyl or aromatic or alkylaromatic).
 CC T is pref. an octanoyl group.
 CC W is L or hydrogen.
 CC The peptides given in AAK45054 8% are examples of X.
 CC the N terminal substit. peptides and proteins have increased
 CC biological activity as compared with unsubst. peptides or proteins
 CC or peptides subst. at the N terminal with an acetyl gp.
 CC They can be used as antimicrobial agents, antiviral agents,
 CC antitumor agents, antiparasitic agents or spermicides and
 CC can also exhibit other biological functions. They can also be
 CC used in promoting or stimulating wound healing, for the treatment
 CC of external burn, and external and/or internal skin and burn
 CC infections or eye infections.
 CC (Updated on 25 MAR 2003 to correct PN field.)
 XX
 SQ Sequence 21 AA;

Quality Match 77.9%; Score 51; Len 14; Length 21;
 Best Local Similarity 85.7%; Pref. No. 0.25;
 Matches 12: Conservation 1: Mismatches 1: Indels 0: Gaps 0:
 27 2 AKALAKALKAL 15
 10 AAK45054 standard: peptide: 21 AA.
 XX
 AC AAK45054;
 XX
 DT 25-MAR-2003 (updated)
 DT 26 JUN 1994 (first entry)
 XX
 XX Basic (positively charged) polypeptide for N terminal
 DE lipophilic substitution.
 KW ion channel; madarin; RNA; XRP; GTP; coenzyme; sarcosine;
 KW amphiphilic; hydrophobic; hydrophilic; lipophilic; growth;
 KW inhibition; target cell; virus; virally-infected cell;
 KW antitumour; antiviral; antitumor; antiparasitic;
 KW spermicide; wound healing; burn; infection.
 XX
 OS Synthetic.
 XX
 ON W09424138-A1.
 XX
 PD 09-DEC-1993.
 XX
 PF 27 MAY 1993; 94W0 US05192.
 XX
 PR 01 JUN 1993; 92W0 US91201.
 XX
 PA (MAGA) MAGAININ PHARM INC.
 XX
 PI Karl B;
 XX
 PR 1993-405419/50.
 XX
 XX Peptide(s) or proteins with an N-terminal lipophilic substit. -
 PI used for inhibiting growth of target cell, virus or
 PI virally-infected cell
 XX
 PS Disclosure: Page 75-88; 113pp; English.
 XX
 CC A novel compsn. for inhibiting growth of a target cell, virus or
 CC virally-infected cell comprises a peptide of formula T-N(W)-X (1),
 CC X is a biologically active amphiphilic ion channel-forming peptide
 CC or protein; pref. a magainin peptide, a pGLa peptide, a XRP
 CC peptide, a CPP peptide, a coetropin or a sarcotoxin.
 CC N is the nitrogen of the N-terminal amino group.
 CC T is a lipophilic moiety; pref. R-50, where R is a 2-hex
 CC hydrocarbon (alkyl or aromatic or alkylaromatic).
 CC T is pref. an octanoyl group.
 CC W is L or hydrogen.
 CC The peptides given in AAK45054 8% are examples of X.
 CC the N terminal substit. peptides and proteins have increased
 CC biological activity as compared with unsubst. peptides or proteins
 CC or peptides subst. at the N terminal with an acetyl gp.
 CC They can be used as antimicrobial agents, antiviral agents,
 CC antitumor agents, antiparasitic agents or spermicides and
 CC can also exhibit other biological functions. They can also be
 CC used in promoting or stimulating wound healing, for the treatment
 CC of external burn, and external and/or internal skin and burn
 CC infections or eye infections.
 CC (Updated on 25 MAR 2003 to correct PN field.)
 XX
 SQ Sequence 21 AA;

Matches 12: Conservation 1: Mismatches 1: Indels 0: Gaps 0:
 27 2 AKALAKALKAL 15
 10 AAK45054 standard: peptide: 21 AA.
 XX
 AC AAK45054;
 XX
 DT 25-MAR-2003 (updated)
 DT 26 JUN 1994 (first entry)
 XX
 XX Basic (positively charged) polypeptide for N terminal
 DE lipophilic substitution.
 KW ion channel; madarin; RNA; XRP; GTP; coenzyme; sarcosine;
 KW amphiphilic; hydrophobic; hydrophilic; lipophilic; growth;
 KW inhibition; target cell; virus; virally-infected cell;
 KW antitumour; antiviral; antitumor; antiparasitic;
 KW spermicide; wound healing; burn; infection.
 XX
 OS Synthetic.
 XX
 ON W09307892-A1.
 XX
 PR 29-APR-1993.
 XX
 PD 16-OCT-1992; 92W0 US08824.
 XX
 PF 16-OCT-1991; 91US 0778771.
 XX
 PA (CHILL) CHILLRENS HOSPITAL PHILADELPHIA.
 XX
 PI Berkowitz B; Eastaff M;
 XX
 PR 1993-162194/18.
 XX
 XX Inhibiting growth of bacteria - by co-administering an anti-
 PI antibiotic and ion channel-forming peptide, even, mutation
 XX
 PS Disclosure: Page 26; 125pp; English.
 XX
 CC The sequence is that of a basic polypeptide of at least 8 hydrophobic
 CC amino acids and at least 8 hydrophilic amino acids. The peptide is
 CC amphiphilic, positively charged and ion channel-forming and may
 CC be used in a compsn. with an antibiotic which is not an X channel
 CC forming peptide, to inhibit the growth of target cells. The peptide
 CC is pref. a magainin peptide, XRP, pGLa or CPP peptide. The peptide
 CC and antibiotic potentiate each other, i.e., interaction of the
 CC peptide with the membranes of bacterial cells facilitates
 CC penetration of the cells by the antibiotic. The various agents
 CC requires less antibiotic and may have effect at a lower concn. to
 CC antibiotic alone. Apart from the above uses the compound may be
 CC used as preservatives or sterilants.
 CC See also AAK45054 800.
 CC (Updated on 25 MAR 2003 to correct PN field.)
 XX
 SQ Sequence 21 AA;

Quality Match 77.9%; Score 51; Len 14; Length 21;
 Best Local Similarity 85.7%; Pref. No. 0.25;
 Matches 12: Conservation 1: Mismatches 1: Indels 0: Gaps 0:
 27 2 AKALAKALKAL 15
 10 AAK45054 standard: peptide: 21 AA.
 XX
 AC AAK45054;
 XX
 DT 25-MAR-2003 (updated)
 DT 26 JUN 1994 (first entry)
 XX
 XX Basic (positively charged) polypeptide for N terminal
 DE lipophilic substitution.
 KW ion channel; madarin; RNA; XRP; GTP; coenzyme; sarcosine;
 KW amphiphilic; hydrophobic; hydrophilic; lipophilic; growth;
 KW inhibition; target cell; virus; virally-infected cell;
 KW antitumour; antiviral; antitumor; antiparasitic;
 KW spermicide; wound healing; burn; infection.
 XX
 OS Synthetic.
 XX
 ON W09307892-A1.
 XX
 PR 29-APR-1993.
 XX
 PD 16-OCT-1992; 92W0 US08824.
 XX
 PF 16-OCT-1991; 91US 0778771.
 XX
 PA (CHILL) CHILLRENS HOSPITAL PHILADELPHIA.
 XX
 PI Berkowitz B; Eastaff M;
 XX
 PR 1993-162194/18.
 XX
 XX Inhibiting growth of bacteria - by co-administering an anti-
 PI antibiotic and ion channel-forming peptide, even, mutation
 XX
 PS Disclosure: Page 26; 125pp; English.
 XX
 CC The sequence is that of a basic polypeptide of at least 8 hydrophobic
 CC amino acids and at least 8 hydrophilic amino acids. The peptide is
 CC amphiphilic, positively charged and ion channel-forming and may
 CC be used in a compsn. with an antibiotic which is not an X channel
 CC forming peptide, to inhibit the growth of target cells. The peptide
 CC is pref. a magainin peptide, XRP, pGLa or CPP peptide. The peptide
 CC and antibiotic potentiate each other, i.e., interaction of the
 CC peptide with the membranes of bacterial cells facilitates
 CC penetration of the cells by the antibiotic. The various agents
 CC requires less antibiotic and may have effect at a lower concn. to
 CC antibiotic alone. Apart from the above uses the compound may be
 CC used as preservatives or sterilants.
 CC See also AAK45054 800.
 CC (Updated on 25 MAR 2003 to correct PN field.)
 XX
 SQ Sequence 21 AA;

```

XX 25-MAR-2003 (updated)
DE 10 MAY-1993 (first entry)
XX C-terminal substd. amphiphilic peptide #12.
XX ion-channel forming; ionophore; antibiotic; anti-tumour; anti virus;
KW wound healing.
XX Synthetic.
XX Key Location/Qualifiers
FT Region 1..7
FT Modified site 21
FT /label= repeat_unit
FT /note= "Leu-(C O) T,
FT T O R, Eit RH2, NH OH or SR'R",
FT R opt.substd. 1-10C aliphatic, aromatic or
FT alkyl ap.;
FT R', R" = H or from one of ops. i and ii;
FT qp.i 1-10C hydroxy-substd. aliphatic,
FT aromatic or alkyl ap.;
FT qp.ii amino-substd. aliphatic, aromatic,
FT alkyl or alkylaromatic qp. and
FT at least one of R' and R" qp.i or qp.ii"
XX
XX W09222417-A1.
XX
XX 23-DEC-1992.
XX
XX 01 JUN-1992: 92W0-US04603.
XX
XX 12 JUN-1993: 91US-0713716.
XX
XX (MAGNA) MAGNIN PHARM INC.
XX
XX Kari UP, Maloy WL;
XX
XX WPI: 1993-017904/02.
XX
XX Now C-terminal-substd. amphiphilic peptide(s) - for treating
XX bacterial, viral or fungal infections and tumours, also useful as
XX spermicide
XX
XX Claim 10: Page 76; 124pp; English.
XX
XX This peptide is a preferred example of a highly generic amphiphilic
XX peptide with a C-terminal modification which increases the peptide's
XX biological activity i.e. the unmodified peptide. The preferred
XX C-terminal modification is -(60)-NHCH2CH2OH or -(20)-NHCH2CH2NH2.
XX Such substd. peptides may be used for inhibiting the growth of a
XX target cell, virus or virally infected cell in a host. The peptides
XX have a broad range of potent antibiotic activity, e.g. against gram-
XX negative and gram positive bacteria, fungi, protozoa and parasites.
XX The peptides can also be used to promote wound healing and treatment
XX of burns, other preferred amphiphilic peptides include maduinins and
XX their analogues, pBA, XPE, cpe, a over-ops and a sarcolexin.
XX
XX (Updated on 25-MAR-2003 to correct IN field.)
XX
XX Sequence 21 AA:
XX
XX Query Match 77.8%; Score 51; DB 14; Length 21;
XX Best Local Similarity 86.7%; Pref. Res. 0.25;
XX Matches 12; Conservative 1; Mismatches 1; Gaps 0;
XX
XX 2 AAALKALLEALKAL 15
XX ||||| |||||
XX 4 SRALKALSKALKAL 17
XX
XX RESULT 7
XX AAR45301
XX ID AAR45301 standard; peptide; 21 AA.

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XX AAR45301;
XX
XX 25 MAR 2003 (updated)
DE 07-JUN-1993 (first entry)
XX Amphiphilic peptide #42 used to treat oral infections.
XX
XX Adverse oral conditions; amphipathic; anti bacterial; anti viral;
XX anti fungal; dental plaque; dental caries; periodontal disease;
XX gingivitis; ionophore; ion channel forming.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FT Region 1..7
FT /note= "repeat unit"
XX
XX W09301723-A1.
XX
XX 04 FEB 1993.
XX
XX 09 JUL-1992: 92W0-US05411.
XX
XX 25 JUL-1993: 91US 0745070.
XX
XX (MAGNA) MAGNIN PHARM INC.
XX
XX Berkowitz B, Jacob LJ
XX
XX WPI: 1993-078434/07.
XX
XX Peptide(s) for prophylaxis and treatment of oral disorders caused
XX for periodontal disease, plaque, dental caries, gingivitis etc.
XX
XX Claim 2: Page 42; 144pp; English.
XX
XX This is a specific example of a highly generic formula of a
XX preferred amphiphilic peptides for use in prevention or treatment of
XX adverse oral conditions. The peptide is an octapeptide (1-8 amino acids)
XX channel forming peptide which has anti bacterial and anti viral activity
XX fungal activity etc, making it suitable for use as an expectorant
XX to treat or prevent periodontal disease, plaque, dental caries,
XX halitosis and gingivitis. The anti bacterial activity will make the
XX useful against bacteria associated with dental caries and gingivitis, and
XX the peptides can stimulate the healing of wounds in the oral cavity.
XX
XX (Updated on 25-MAR-2003 to correct IN field.)
XX
XX Sequence 21 AA:
XX
XX Query Match 77.8%; Score 51; DB 14; Length 21;
XX Best Local Similarity 86.7%; Pref. Res. 0.25;
XX Matches 12; Conservative 1; Mismatches 1; Gaps 0;
XX
XX 2 AAALKALLEALKAL 15
XX ||||| |||||
XX 4 SRALKALSKALKAL 17
XX
XX RESULT 8
XX AAR45015
XX ID AAR45015 standard; peptide; 21 AA.
XX
XX AAR45015;
XX
XX 25 MAR 2003 (updated)
DE 08 MAY 1993 (first entry)
XX
XX Biologically active amphiphilic peptide.
XX
XX Synectostic and imbricoidal composition; compound; the compound is a
XX combination; preservation; structural; functional; activity;
XX control; treatment; prevention; external barrier; etc. etc. etc.

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XX OS Synthetic.
XX
XX FI Inactivation/Qualifiers
XX Modified-site 21
XX
XX FA Inactive "inhibitor"
XX
XX PN W09411783-A1.
XX
XX PD 24-JUN-1994.
XX
XX PF 03-DEC-1992: 92W0-US10427.
XX
XX PR 09-DEC-1991: 910S-0804629.
XX
XX PA (MMAA) MALA-KIN PEAKM IN.
XX
XX PI Belkowitz B.
XX
XX DK WPI: 1993-21-0816/26.
XX
XX PI Synergistic antimicrobial compound, for treatment or prevention
XX of infection of burns - contd. amphiphilic peptide or protein e.g.
XX maintain peptide and chelating agent
XX
XX PS Example: Page 24: 122pp: Endlish.
XX
XX CC The sequence is that of a biologically active amphiphilic peptide
XX which is used in a synergistic antimicrobial compound with a
XX chelating agent. The chelating agent potentiates the activity of the
XX peptide by binding inhibitory Ca/Mg ions and may also increase
XX permeability of the target cells to the peptide. The compound is
XX active against a wide range of microorganisms (esp. bacterial) and
XX can be used as a preservative or sterilant, or to control infections
XX in animals or plants. Particular applications are to treat/prevent
XX infections of external burns and treatment of eye infections e.g.
XX where the pathogen is Pseudomonas aeruginosa, Staphylococcus aureus,
XX Streptococcus or Neisseria gonorrhoeae.
XX (Updated on 25-MAR-2003 to correct FN field.)
XX
XX Sequence 21 AA:
XX
XX Query Match 77.4% Score 51: 16-14: Length 21:
XX Best Local Similarity 85.7% Prod. No. 0.25:
XX Matches 12: Conservative 1: Mismatches 1: Indels 0: Gaps 0:
XX
XX QY 2 AKAKKALKKALKAL 15
XX
XX DI 4 SKAKKALKKALKAL 17
XX
XX DE
XX
XX RESULT 9
XX AAR55907
XX ID AAR55907 standard: peptide: 21 AA.
XX
XX AV AAR55907:
XX
XX DI 25-MAR-2003 (updated)
XX 16-DEC-1994 (first entry)
XX
XX XX
XX DE Amphiphilic peptide #42.
XX
XX KW Amphiphilic: ion forming; dyanocolorical malianney; polyanion;
XX XPP; cyp; cypoxin; modifing; apidoxin; apidoxin;
XX KW major basic protein; eosinophilic; other; cypoxin;
XX KW bacterial permeability increasing protein; cypoxin;
XX
XX OS Synthetic.
XX
XX IN W09411783-A1.
XX
XX PD 17-MAR-1994.
XX
XX PF 16-AUG-1993: 92W0-US07798.
XX
XX PR 31-AUG-1992: 920S-0907462.
XX
XX FA (MMAA) MALA-KIN PEAKM IN.
XX
XX PI Baker MA. JACS 125: Mar 9: 1903.
XX
XX PD WPI: 1994-10085/12.
XX
XX CC Treating dyanocolorical tumours with amphiphilic peptide(s)
XX which form ion channels, esp. maintain cell membrane
XX PD failure, for treating dyanocolorical tumours with amphiphilic
XX

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```

XX
XX PF 03-DEC-1993: 92W0-US11885.
XX
XX PR 03-DEC-1992: 920S-0984957.
XX
XX FA (MMAA) MALA-KIN PEAKM IN.
XX
XX PI Jacob LS. Maloy WL.
XX
XX DR WPI: 1994-199963/24.
XX
XX PI Treating skin cancer with ion channel forming peptide(s) - e.g.
XX malianney, modifing etc., specifically for treating melanoma
XX
XX PS Disclature: Page 84: 12pp: Endlish.
XX
XX CC The peptide is used to treat dermatological melanocytic
XX may be used to treat especially melanoma but also basal, squamous
XX squamous cell carcinomas. It can be used together with an anti-
XX also inhibits/greets growth of the target cells. Peptides used for
XX such therapy include malianney, 16-14 or cyp peptide, cypoxin,
XX sarcosine, melittin, apidoxin, defensins, etc. (first entry)
XX cypoxin; cypoxin; bactericidal permeability increasing protein
XX See also AAR55907: 05597.
XX (Updated on 25-MAR-2003 to correct FN field.)
XX
XX Sequence 21 AA:
XX
XX Query Match 77.4% Score 51: 16-14: Length 21:
XX Best Local Similarity 85.7% Prod. No. 0.25:
XX Matches 12: Conservative 1: Mismatches 1: Indels 0: Gaps 0:
XX
XX QY 2 AKAKKALKKALKAL 15
XX
XX DI 4 SKAKKALKKALKAL 17
XX
XX DE
XX
XX RESULT 10
XX AAR55904
XX ID AAR55904 standard: peptide: 21 AA.
XX
XX AV AAR55904:
XX
XX DI 25-MAR-2003 (updated)
XX 16-DEC-1994 (first entry)
XX
XX XX
XX DE Amphiphilic peptide #42.
XX
XX KW Amphiphilic: ion forming; dyanocolorical malianney; polyanion;
XX XPP; cyp; cypoxin; modifing; apidoxin; apidoxin;
XX KW major basic protein; eosinophilic; other; cypoxin;
XX KW bacterial permeability increasing protein; cypoxin;
XX
XX OS Synthetic.
XX
XX IN W09411783-A1.
XX
XX PD 17-MAR-1994.
XX
XX PF 16-AUG-1993: 92W0-US07798.
XX
XX PR 31-AUG-1992: 920S-0907462.
XX
XX FA (MMAA) MALA-KIN PEAKM IN.
XX
XX PI Baker MA. JACS 125: Mar 9: 1903.
XX
XX PD WPI: 1994-10085/12.
XX
XX CC Treating dyanocolorical tumours with amphiphilic peptide(s)
XX which form ion channels, esp. maintain cell membrane
XX PD failure, for treating dyanocolorical tumours with amphiphilic
XX

```


AAR90075
 ID AAR90075 standard; peptide; 21 AA.
 XX
 AC AAR90075;
 XX
 DT 03-JUL-1996 (first entry)
 DT
 XX (KALSKAL) 3 peptide modified by N-terminal lipophilic group.
 DE
 XX Ion channel forming peptide; lipophilic; N-terminal modification;
 KW maddinin; inhibition; cell growth; viral replication; ionophore;
 KW membrane permeability; antimicrobial; anti-bacterial; antibiotic;
 KW anti-fungal; anti-viral; spermicidal; anti-tumour; anti-parasitic.
 XX
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FH Modified-site
 FT /note: "N-terminal amino group is mono- or
 FT di-subst. by lipophilic moiety, esp.
 FT octanoyl"
 FT
 FT Peptide 1..7
 FT /label repeat
 FT /note: "first of 3 repeats of amphipathic
 FT heptapeptide motif"
 FT
 FN W09519470-A1.
 XX
 XX 20-JUL-1995.
 PD
 XX
 XX 18-JAN-1995; 95W0-US00714.
 XX
 XX 18-JAN-1994; 94US-0184462.
 XX
 XX (MAGA-) MAGAININ PHARM INC.
 XX
 DT Kari UP, McLane M, Williams TJ;
 XX
 DR WPI: 1995-263826/34.
 XX
 XX Ion channel-forming amphiphilic peptide(s) with N-terminal
 PT lipophilic gps. - useful e.g. as antiviral, antibacterial,
 PT antiparasitic or antitumour agents
 XX
 PS Claim 21: Page 78: 139pp; English.
 XX
 CC The present peptide is a specific example corresp. to a highly
 CC generic formula for ion channel forming peptides (ionophores)
 CC consisting of 1-5 repeats of an amphipathic heptapeptide motif.
 CC these ionophores are known to have a broad range of potent
 CC antibiotic activity against microorganisms including gram-positive
 CC and gram-negative bacteria, fungi, viruses, protozoa and parasites.
 CC N-terminal modification (pref. mono-substn. by octanoyl) to produce
 CC an ion channel forming peptide having a lipophilic N-terminus
 CC increases the biological activity of the peptides against target cells,
 CC viruses and virally-infected cells, compared to peptides substd. with
 CC an acetyl group at the N-terminus. Compositions comprising the peptides
 CC with lipophilic modifications are claimed for inhibiting growth of a
 CC target cell, virus or virally-infected cell.
 XX
 SQ Sequence 21 AA;
 Query Match 77.3%; Score 51; DB 16; Length 21;
 Best Local Similarity 85.7%; Pred. No. 0.25;
 Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 2 AKALKALKALKAL 15
 DB 4 SKALKALKALKAL 17
 RESULT 14
 AAR93846

AAR93846 standard; peptide; 21 AA.
 XX
 AC AAR93846;
 XX
 DT 25-MAR-2003 (updated)
 DT 24-MAY-1996 (first entry)
 XX
 DE Ion channel forming amphiphilic peptide #6.
 XX
 KW Ion-channel; amphiphilic peptide; antimicrobial; antiviral; antitumor;
 KW antitumor; antiparasitic; antitumoral; spermicide; cyst; spore; skin;
 KW trophozoite; infection; wound healing; burn.
 XX
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FH Misc-difference 21
 FT /note: "contains C-terminal amide group"
 XX
 FN US5459237-A.
 XX
 XX 17 OCT-1995.
 PD
 XX
 XX 02-JUL-1992; 92US-0908455.
 XX
 XX 02-JUL-1992; 92US-0908455.
 PR 08-FEB-1990; 90US-0476629.
 PR 15-APR-1991; 91US-0686115.
 XX
 XX (MAGA-) MAGAININ PHARM INC.
 XX
 DT Berkowitz B, Kari UP, Maloy WL;
 XX
 DR WPI: 1995-365836/47.
 XX
 XX New ion channel-forming amphiphilic peptide(s) useful as
 PT antimicrobial, antiviral, antitumor, antiparasitic or antitumoral
 PT agents or for wound healing
 XX
 PS Disclosure: Column 5: 38pp; English.
 XX
 CC The peptides AAR93841-93808 are examples of novel ion-channel forming
 CC amphiphilic peptides which can be used as antimicrobial, antiviral,
 CC antibiotic, antitumor, antiparasitic, antitumoral agents or as
 CC spermicides. The peptides can be used as preservatives or sterilants
 CC for materials susceptible to microbial or viral infections. The
 CC peptides can also be used for killing cysts, spores or trophozoites of
 CC infection-causing organisms. They can also be used to promote wound
 CC healing or to treat or prevent skin and burn infections. The peptides
 CC can be administered in combination with other biologically active
 CC amphiphilic peptides such as AAR93909-29.
 XX
 XX (Updated on 25-MAR-2003 to correct PF field.)
 XX
 SQ Sequence 21 AA;
 Query Match 77.3%; Score 51; DB 16; Length 21;
 Best Local Similarity 85.7%; Pred. No. 0.25;
 Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 2 AKALKALKALKAL 15
 DB 4 SKALKALKALKAL 17
 RESULT 15
 AAY10671
 ID AAY10671 standard; peptide; 21 AA.
 XX
 AC AAY10671;
 XX
 DT 11-MAY-1999 (first entry)
 DT
 XX Peptide used to make biologically active peptides.

XX Superficial septic shock; bacteraemia; meningitis; cystic fibrosis;
 KW antimicrobial; antiviral; antibacterial; antifungal; antitumour;
 KW antiparasitic; spermicide; preservative; sterility; disinfectant;
 KW wound healing; burn; skin infection; eye infection; solid tumour;
 KW leukaemia; non-small cell lung cancer; adenocarcinoma; plant infection;
 KW periodontal disease; plaque; gingivitis; caries; Streptococcus mutans.
 XX Synthetic.
 OS W090486 A2.
 IN 18 JAN 1994;
 XX 01 JUN 1994;
 PD 28 JAN 1999;
 XX 15 JUL 1998;
 XX 15 JUL 1998;
 PR 15 JUL 1997;
 XX (MAGA-) MACAININ PHARM INC.
 PA Kari DP, Williams TJ, Melane M;
 XX WPI: 1999-141869/11.
 XX
 XX Treating sepsis or septic shock with N-methylated ion-channel forming
 PT peptide - or its methane sulphonate derivative of reduced toxicity.
 PT also generally useful as antimicrobial and antitumour agents.
 XX
 PS Disclosure: Page 154; 202pp; English.
 XX
 XX AAV1040 795 represent peptides used in the production of biologically
 CC active peptides with reduced toxicity. The biologically active peptides
 CC are used to treat sepsis or septic shock, and comprise the formula:
 CC i (N⁺) X, where X is biologically active, amphipathic, ion-channel
 CC forming peptide or protein; 1. Lipophilic group; and 2. Hydroxyl or T.
 CC The peptides are particularly used to treat infections by pseudomonas
 CC aeruginosa in patients with cystic fibrosis, but more generally as
 CC antimicrobial, antiviral, antibacterial, antifungal, antitumour or
 CC antiparasitic agents, and also as spermicides, e.g., as preservatives,
 CC sterilants, and disinfectants in human and veterinary medicine. They
 CC can be used to stimulate wound healing, treat burns and/or skin and
 CC burn infections, eye infections, solid tumours or leukaemia
 CC (particularly non-small cell lung cancer and adenocarcinoma, including
 CC those resistant to other antitumour agents), and also for treatment of
 CC infections in plants, and, when formulated in oral hyaline formulations,
 CC for treating or preventing periodontal disease, plaque, gingivitis and/or
 CC caries (specifically by action on Streptococcus mutans).
 XX
 SQ Sequence: 21 AA;

Query Match: 77.8%; Score 51; DB 20; Length 21;
 Best Local Similarity: 86.7%; Pred. NO: 0.25;
 Matches: 12; Conservative: 1; Mismatches: 1; Indels: 0; Gaps: 0;

CV 2 AKAIKALIKAL 15
 ID 1111111111
 4 SEAKALIKAL 17
 RESULT 16
 AAE22484
 ID AAE22484 standard; peptide: 21 AA.
 XX
 AC AAE22484:
 XX
 DT 25 JUL 2002 (first entry)
 XX
 DE Biologically active peptide #16.
 XX
 KW Biologically active peptide; toxicity; antimicrobial; antitumour;
 KW methane sulphonate derivative; wound healing; burn; therapy; sepsis;
 KW eye infection; cyst; sporotrichosis; tumour; lung infection;
 KW cystic fibrosis; septic shock; bacterial endocarditis; pharyngitis;

KW antibacterial; immunosuppressive;
 XX
 OS Unidentified.
 XX 08648445 R1.
 XX 19 FEB 2002;
 XX 15 JUL 1998;
 XX 15 JUL 1998;
 PR 01 JUN 1994;
 PR 01 JUN 1994;
 PR 05 JUN 1995;
 PR 15 JUL 1997;
 XX (MAGA-) MACAININ PHARM INC.
 PA Kari DP, Williams TJ, Melane M;
 XX WPI: 2002-00079/08.
 XX
 XX Reducing toxicity of unsubstituted or N-terminal substituted peptide
 PT having antimicrobial and antitumor activity useful in treatment of
 PT infections and tumor, by forming methane sulphonate derivative of
 PT peptide.
 XX
 PS Disclosure: Column 17; 16pp; English.
 XX
 XX The invention relates to biologically active peptides which have
 CC toxicity and methods of preparing them. The peptides are generally
 CC used to treat sepsis or septic shock, and comprise the formula:
 CC while exhibiting reduced toxicity. The method of forming the
 CC involves the formation of related methane sulphonate derivative
 CC analogues. The method is useful for reducing toxicity of
 CC unsubstituted peptide or an N-terminal substituted peptide which
 CC utilized in promoting or stimulating healing of a wound, or in
 CC treatment of external burn, prevention or treatment of infection
 CC caused by bacteria or fungi, in killing cysts, e.g., in cystic
 CC of infection causing organisms, and may also be employed in the
 CC treatment of tumors, serious lung infections such as those caused in
 CC in cystic fibrosis, for treating sepsis, septic shock, and other
 CC related ailments, and for neutralising toxicity caused by the
 CC present sequence. It is a biologically active peptide of the invention.
 XX
 SQ Sequence: 21 AA;
 Query Match: 77.8%; Score 51; DB 20; Length 21;
 Best Local Similarity: 86.7%; Pred. NO: 0.25;
 Matches: 12; Conservative: 1; Mismatches: 1; Indels: 0; Gaps: 0;

CV 2 AKAIKALIKAL 15
 ID 1111111111
 4 SEAKALIKAL 17

RESULT 17
 AAE22484
 ID AAE22484 standard; peptide: 14 AA.
 XX
 AC AAE22484:
 XX
 DT 02 OCT 1998 (first entry)
 XX
 DE Minimalist lytic peptide.
 XX
 KW Lytic peptide; channel forming peptide; antitumour; pharyngitis;
 XX
 OS Synthetic.
 XX
 EN US5786542 A.
 XX
 PD 04 AUG 1998.
 XX


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PF 06-OCT-1997; 970S-0944133.
XX
PR 22-APR-1994; 940S-0242525.
PR 22-JUL-1996; 960S-0681075.
PR 03-FEB-1997; 970S-0789077.
PR 06-OCT-1997; 970S-0944133.
XX
PA (LOUIS ) UNIV LOUISIANA STATE & AGRIC & MECH COLL.
XX
XX Becker CL, McLaughlin ML;
XX WPI; 1998-446184/38.
XX
XX Selective lysis of bacteria amongst mammalian cells - using 14-mer
XX or 21-mer lytic peptides
XX
XX claim 7; Column 35; 25pp; English.
XX
XX AAW62920-67 represent minimalist lytic (channel forming) peptides. The
XX peptides have antibacterial properties in concentrations not lethal
XX toward mammalian cells. The peptides are heptads (or heptad multimers)
XX that comprise four nonpolar amino acid residues and three positively
XX charged amino acid residues, or five nonpolar amino acid residues and
XX two positively charged amino acid residues. The nonpolar amino acid
XX residues and the positively charged amino acid residues are distributed
XX within the heptad such that when the multimer forms an alpha-helix the
XX nonpolar amino acid residues will lie on one face of the alpha-helix,
XX and the positively charged amino acid residues will lie on the opposite
XX face of the alpha-helix, whereby the multimer is amphipathic.
XX
SQ Sequence 14 AA;

Query Match 75.8%; Score 50; DB 19; Length 14;
Best Local Similarity 92.3%; Pred. No. 0.24;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 KALKALKALKAL 15
DB 2 KALKALKALKAL 14

RESULT 18
AAW62933
XX
XX AAW62933 standard; peptide; 14 AA.
XX
XX AAW62933;
XX
XX 02-OCT-1998 (first entry)
XX
XX Minimalist lytic peptide.
XX
XX Lytic peptide; channel forming peptide; antibacterial; amphipathic.
XX
XX Synthetic.
XX
XX US5789542-A.
XX
XX 04-AUG-1998.
XX
XX 06-OCT-1997; 970S-0944133.
XX
XX 22-APR-1994; 940S-0242525.
XX 22-JUL-1996; 960S-0681075.
XX 03-FEB-1997; 970S-0789077.
XX 06-OCT-1997; 970S-0944133.
XX
XX (LOUIS ) UNIV LOUISIANA STATE & AGRIC & MECH COLL.
XX
XX Becker CL, McLaughlin ML;
XX WPI; 1998-446184/38.
XX
XX Selective lysis of bacteria amongst mammalian cells - using 14-mer

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PF or 21-mer lytic peptides
XX
XX claim 6; Column 35; 25pp; English.
XX
XX AAW62920-67 represent minimalist lytic (channel forming) peptides. The
XX peptides have antibacterial properties in concentrations not lethal
XX toward mammalian cells. The peptides are heptads (or heptad multimers)
XX that comprise four nonpolar amino acid residues and three positively
XX charged amino acid residues, or five nonpolar amino acid residues and
XX two positively charged amino acid residues. The nonpolar amino acid
XX residues and the positively charged amino acid residues are distributed
XX within the heptad such that when the multimer forms an alpha-helix the
XX nonpolar amino acid residues will lie on one face of the alpha-helix,
XX and the positively charged amino acid residues will lie on the opposite
XX face of the alpha-helix, whereby the multimer is amphipathic.
XX
SQ Sequence 14 AA;

Query Match 75.8%; Score 50; DB 19; Length 14;
Best Local Similarity 92.3%; Pred. No. 0.24;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 KALKALKALKAL 15
DB 1 KALKALKALKAL 13

RESULT 19
AAW62946
XX
XX AAW62946 standard; peptide; 21 AA.
XX
XX AAW62946;
XX
XX 02-OCT-1998 (first entry)
XX
XX Minimalist lytic peptide.
XX
XX Lytic peptide; channel forming peptide; antibacterial; amphipathic.
XX
XX Synthetic.
XX
XX US5789542-A.
XX
XX 04-AUG-1998.
XX
XX 06-OCT-1997; 970S-0944133.
XX
XX 22-APR-1994; 940S-0242525.
XX 22-JUL-1996; 960S-0681075.
XX 03-FEB-1997; 970S-0789077.
XX 06-OCT-1997; 970S-0944133.
XX
XX (LOUIS ) UNIV LOUISIANA STATE & AGRIC & MECH COLL.
XX
XX Becker CL, McLaughlin ML;
XX WPI; 1998-446184/38.
XX
XX Selective lysis of bacteria amongst mammalian cells - using 14-mer
XX or 21-mer lytic peptides
XX
XX Disclosure; Column 6; 25pp; English.
XX
XX AAW62920-67 represent minimalist lytic (channel forming) peptides. The
XX peptides have antibacterial properties in concentrations not lethal
XX toward mammalian cells. The peptides are heptads (or heptad multimers)
XX that comprise four nonpolar amino acid residues and three positively
XX charged amino acid residues, or five nonpolar amino acid residues and
XX two positively charged amino acid residues. The nonpolar amino acid
XX residues and the positively charged amino acid residues are distributed
XX within the heptad such that when the multimer forms an alpha-helix the
XX nonpolar amino acid residues will lie on one face of the alpha-helix,
XX and the positively charged amino acid residues will lie on the opposite

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XX FN W09012866 A1.
 XX ID 01 NOV 1990.
 XX PE 10 APR 1990; 90W0-US0445.
 XX PR 10 APR 1989; 89US-036181.
 XX PA (DOMINION) UNIVERSITY OF LOUISIANA STATE.
 XX JAYNES JM.
 XX WPI: 1990 48846/46.
 XX New lytic polypeptide(s) with proliferative activity - are
 alpha helical peptide(s) having aligned amphipathic for treating
 microbial infections and lysing cancer cells
 Claim 14; page 40; 57pp; English.
 CC This peptide is an analogue of a known lytic peptide. It comprises
 an alpha helical conformation of amino acids. It is effective at
 lysing e.g. gram-positive and -negative bacteria and mammalian neo-
 plastic cells, cells infected with intracellular pathogenic micro-
 organisms such as HIV. It stimulates the proliferation of fibro-
 blasts and lymphocytes and can be used in wound healing.
 See also AAR07744-47, AAR07739-41 and AAR07743-51.
 CC (Updated on 25-MAR-2003 to correct PA field.)
 XX Sequence 24 AA:
 QUOTY Match 72.7%; Score 48; DB 11; Length 24;
 Best Local Similarity 80.0%; Prod. No. 0.8;
 Matches 12; Conservative 0; Mismatches 0; Gaps 0;
 QY 1 FAKALKALKALKAL 15
 DB 11 1111 1111 1
 DB 1 FALAKALKALKALKAL 15
 RESULT 24
 ID AAR84166 standard; peptide: 24 AA.
 XX AAR84166;
 XX DT 06-JUN-1996 (first entry)
 XX DE Peptide enhance of fibroblast and keratinocyte proliferation.
 XX amphipathic peptide; enhance; fibroblast; keratinocyte; proliferation;
 KW wound healing; defensin; antimicrobial.
 XX US Synthetic.
 XX W09528832 A1.
 XX 02-NOV 1995.
 XX 19 APR 1995; 95W0-US04718.
 XX 20-APR 1994; 94US-0241730.
 XX (DEME) DEMETER BIO-TECHNOLOGIES LTD.
 XX JAYNES JM. Julian GR.
 XX WPI: 1995 482791/49.
 XX Use of amphipathic peptide(s) for enhancing fibroblast and
 keratinocyte proliferation to promote wound healing in a mammal

PS Claim 7; Page 55; 64pp; English.
 XX AAR84128-73 are amphipathic peptides which are able to stimulate the
 CC proliferative growth of fibroblasts and epithelial cells such as
 CC keratinocytes, hence enhance wound healing in mammalian skin wounds.
 CC The peptides concomitantly have an antimicrobial effect against
 CC microbial species including those which cause of other disease mediators
 CC sepsis and wound infection.
 XX Sequence 24 AA:
 QUOTY Match 72.7%; Score 48; DB 11; Length 24;
 Best Local Similarity 80.0%; Prod. No. 0.8;
 Matches 12; Conservative 0; Mismatches 0; Gaps 0;
 QY 1 FAKALKALKALKAL 15
 DB 11 1111 1111 1
 DB 1 FALAKALKALKALKAL 15
 RESULT 24
 ID AAR77080 standard; peptide: 24 AA.
 XX AAR77080;
 XX DT 20-MAY 1996 (first entry)
 XX DE Synthetic anti neoplastic lytic peptide.
 XX Anti cancer; lysis; amphipathic; neoplasia; cancer; cystic fibrosis;
 KW bronchopulmonary; viral; virus; anti cancer; anti neoplastic;
 KW melittin; defensin.
 XX US Synthetic.
 XX W09527497 A1.
 XX 19-OCT 1995.
 XX 06-APR 1995; 95W0-US04435.
 XX 08-APR 1994; 94US-0225476.
 XX (DEME) DEMETER BIO-TECHNOLOGIES LTD.
 XX JAYNES JM. Julian GR.
 XX WPI: 1995 466226/47.
 XX Method of combatting mammalian neoplasias and other disease states
 PT by delivering non-naturally occurring, non-immunogenic, anti-cancer
 PT proliferative lytic peptide
 PS Claim 7; Page 53; 64pp; English.
 XX AAR77042-R77081 are synthetic, amphipathic, lytic peptides and analogues
 CC melittin, cecropin, manniin and defensin peptides. The peptides are
 CC between 24 and 49 residues long, are amphipathic, carry an overall
 CC positive charge and have anti neoplastic activity. The peptides are
 CC specifically useful for the lysis of cancer cells. Mammalian
 CC cells are resistant to lysis due to their highly organised
 CC cytoskeleton, cancerous cells however possess an altered and
 CC structurally compromised cytoskeleton which when acted upon by lytic
 CC peptides will cause cell lysis. Thus as the lytic peptides to be
 CC used for in vivo treatment of cancers, the peptides are especially
 CC for the treatment of female mammalian cancers such as breast, ovarian,
 CC uterine and cervical cancers. The peptides can however be used for
 CC treat most forms of cancer, cystic fibrosis, bronchopulmonary
 CC and bronchopulmonary viral and microbial infections.

XX Sequence 24 AA;

```

Query Match          ZZ:78; Score 48; DB 16; Length 23;
Best Local Similarity 80.0%; Pred. No. 0.8;
Matches 12; Conservat 100; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKL 15

RESULT 25
AA092415
ID AAR92415 standard; peptide: 23 AA.
XX
AC AAR92415;
GI 17 SEP-1996. (first entry)
XX
DE Lytic peptide used in ubiquitin-lytic peptide fusion protein.
XX
KW Ubiquitin fusion protein; lysis; infection; neoplasia; wound healing;
KW stability; reduced toxicity.
XX
OS Synthetic.
FN W09603519 AL.
XX
PD 08-FEB-1996.
XX
XX 24 JUL 1995; 95WO-0509349.
XX
XX 22 JUL 1994; 94DS-0279472.
XX
XX (DEME-) DEMETER BIOTECHNOLOGIES LTD.
XX (US) } IN SP. OF ASPEC.
XX
XX Belknap W, Garbarino J, Jaynes J;
XX
XX WPI; 1996 117061/12.
XX
XX New fusion protein of ubiquitin and a lytic peptide - for treating
XX infections and neoplasia, healing wounds, etc., also related nucleic
XX acid, vectors, and transformed cells
XX
XX Claim 5; Page 23; 11pp; English.
XX
XX AAR92415 R92462 are lytic peptides used to create ubiquitin-lytic
XX peptide fusion proteins in which the ubiquitin polypeptide is linked
XX at its C' terminus to the lytic peptide. The lytic peptides are pre-
XX selected from either the ceeropins, defensins, sarcotoxins, melittin
XX and maitins. The fusion proteins (FPS) are useful for treating
XX protozoal, bacterial, fungal and viral infections and neoplasia (in
XX plants and animals) in the same way as the FP alone, they also
XX promote wound healing. FPS produced in bacteria may be cleaved in
XX vitro by ubiquitin hydrolases to recover the active lytic peptide.
XX FPS produced in eukaryotic cells are cleaved by endogenous enzymes
XX to yield lytic peptide. Recombinant DNA encoding the FPS have
XX greater stability in bacteria than DNA encoding the lytic peptide
XX only.
XX
XX Sequence 23 AA;

Query Match          ZZ:78; Score 48; DB 17; Length 23;
Best Local Similarity 80.0%; Pred. No. 0.8;
Matches 12; Conservat 100; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKL 15

RESULT 25
AA092415
ID AAR92415 standard; peptide: 23 AA.
XX
AC AAR92415;
GI 17 SEP-1996. (first entry)
XX
DE Lytic peptide used in ubiquitin-lytic peptide fusion protein.
XX
KW Ubiquitin fusion protein; lysis; infection; neoplasia; wound healing;
KW stability; reduced toxicity.
XX
OS Synthetic.
FN W09603519 AL.
XX
PD 08-FEB-1996.
XX
XX 24 JUL 1995; 95WO-0509349.
XX
XX 22 JUL 1994; 94DS-0279472.
XX
XX (DEME-) DEMETER BIOTECHNOLOGIES LTD.
XX (US) } IN SP. OF ASPEC.
XX
XX Belknap W, Garbarino J, Jaynes J;
XX
XX WPI; 1996 117061/12.
XX
XX New fusion protein of ubiquitin and a lytic peptide - for treating
XX infections and neoplasia, healing wounds, etc., also related nucleic
XX acid, vectors, and transformed cells
XX
XX Claim 5; Page 23; 11pp; English.
XX
XX AAR92415 R92462 are lytic peptides used to create ubiquitin-lytic
XX peptide fusion proteins in which the ubiquitin polypeptide is linked
XX at its C' terminus to the lytic peptide. The lytic peptides are pre-
XX selected from either the ceeropins, defensins, sarcotoxins, melittin
XX and maitins. The fusion proteins (FPS) are useful for treating
XX protozoal, bacterial, fungal and viral infections and neoplasia (in
XX plants and animals) in the same way as the FP alone, they also
XX promote wound healing. FPS produced in bacteria may be cleaved in
XX vitro by ubiquitin hydrolases to recover the active lytic peptide.
XX FPS produced in eukaryotic cells are cleaved by endogenous enzymes
XX to yield lytic peptide. Recombinant DNA encoding the FPS have
XX greater stability in bacteria than DNA encoding the lytic peptide
XX only.
XX
XX Sequence 23 AA;

Query Match          ZZ:78; Score 48; DB 17; Length 23;
Best Local Similarity 80.0%; Pred. No. 0.8;
Matches 12; Conservat 100; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKL 15

RESULT 25
AA092415
ID AAR92415 standard; peptide: 23 AA.
XX
AC AAR92415;
GI 17 SEP-1996. (first entry)
XX
DE Lytic peptide used in ubiquitin-lytic peptide fusion protein.
XX
KW Ubiquitin fusion protein; lysis; infection; neoplasia; wound healing;
KW stability; reduced toxicity.
XX
OS Synthetic.
FN W09603519 AL.
XX
PD 08-FEB-1996.
XX
XX 24 JUL 1995; 95WO-0509349.
XX
XX 22 JUL 1994; 94DS-0279472.
XX
XX (DEME-) DEMETER BIOTECHNOLOGIES LTD.
XX (US) } IN SP. OF ASPEC.
XX
XX Belknap W, Garbarino J, Jaynes J;
XX
XX WPI; 1996 117061/12.
XX
XX New fusion protein of ubiquitin and a lytic peptide - for treating
XX infections and neoplasia, healing wounds, etc., also related nucleic
XX acid, vectors, and transformed cells
XX
XX Claim 5; Page 23; 11pp; English.
XX
XX AAR92415 R92462 are lytic peptides used to create ubiquitin-lytic
XX peptide fusion proteins in which the ubiquitin polypeptide is linked
XX at its C' terminus to the lytic peptide. The lytic peptides are pre-
XX selected from either the ceeropins, defensins, sarcotoxins, melittin
XX and maitins. The fusion proteins (FPS) are useful for treating
XX protozoal, bacterial, fungal and viral infections and neoplasia (in
XX plants and animals) in the same way as the FP alone, they also
XX promote wound healing. FPS produced in bacteria may be cleaved in
XX vitro by ubiquitin hydrolases to recover the active lytic peptide.
XX FPS produced in eukaryotic cells are cleaved by endogenous enzymes
XX to yield lytic peptide. Recombinant DNA encoding the FPS have
XX greater stability in bacteria than DNA encoding the lytic peptide
XX only.
XX
XX Sequence 23 AA;

Query Match          ZZ:78; Score 48; DB 17; Length 23;
Best Local Similarity 80.0%; Pred. No. 0.8;
Matches 12; Conservat 100; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKL 15

RESULT 27
AA076374
ID AAR76374 standard; peptide: 23 AA.
XX
AC AAR76374;
GI 04 DEC 1999. (first entry)
XX
DE Rat leucine peptide.
XX
XX

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XX AAR9972;
XX
XX 16 SEP 1996. (first entry)
XX
XX Synthetic lytic peptide #6.
XX
XX Lytic peptide; ubiquitin; synthetic analogue; cell membrane fusion protein;
XX microbial pathogen; disease resistant plant; bactericidal; cytotoxic; fungicidal;
XX protozoa; virus; neoplasia; fusion protein; hydrolysis.
XX
XX Synthetic.
XX
XX W09603522 AL.
XX
XX 08 FEB 1996.
XX
XX 24 JUL 1995; 95WO-0509349.
XX
XX 22 JUL 1994; 94DS-0279472.
XX
XX (DEME-) DEMETER BIOTECHNOLOGIES LTD.
XX (US) } IN SP. OF ASPEC.
XX
XX Jaynes J;
XX
XX WPI; 1996 117064/12.
XX
XX Lytic peptide(S), useful for developing disease resistant plants,
XX can be expressed as fusion protein with ubiquitin for greater prodn.
XX in bacterial host cells
XX
XX Claim 1; Page 71; 11pp; English.
XX
XX AAR9994749921 and AAR9926 R99763 represent synthetic analogues of
XX naturally occurring lytic peptides. Lytic peptides best known for their
XX and other non-host cells by disrupting the cell membrane and promoting
XX cell lysis. Synthetic lytic peptide analogues have similar or higher
XX levels of lytic activity for many different types of cells compared to
XX naturally occurring forms. The concentration of the synthetic analogues
XX required to lyse microbial pathogens does not vary from natural peptides
XX cells. The lytic peptides can be expressed in plants to develop for
XX development of disease resistant plants. The peptides are useful for
XX promoting wound healing and combating bacterial infections in plants.
XX The lytic peptides can also be used for wound healing in humans.
XX viral or bacterial infections of neoplasia in mammals and plants.
XX Lytic peptide ubiquitin fusion proteins are created by fusing the
XX bacterial hosts. Bacteria lack the hydrolytic activity of the peptide
XX from ubiquitin, and therefore the active (and lytic) peptides peptide
XX will not be released in the host cells. The resultant fusion products
XX lytic peptide can be retrieved from the host cells by a procedure
XX vitro.
XX
XX Sequence 23 AA;

Query Match          ZZ:78; Score 48; DB 17; Length 23;
Best Local Similarity 80.0%; Pred. No. 0.8;
Matches 12; Conservat 100; Mismatches 0; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
DB 1 FALKALKALKALKL 15

RESULT 27
AA076374
ID AAR76374 standard; peptide: 23 AA.
XX
AC AAR76374;
GI 04 DEC 1999. (first entry)
XX
DE Rat leucine peptide.
XX
XX

```

KW GnRH: lytic peptide; rat; gonadotrophin-releasing hormone; contraception;
 KW sterility; uterine hormone; ad; LH-RH-III; fertility; pituitary;
 KW lampry; luteinising hormone-releasing hormone; gonadotropin; inhibitor;
 KW tumour; ligand; fusion protein; reproductive capacity; insect; treatment;
 KW growth rate; pest control; L-hecate; cancer; adenoma; lymphocyte;
 KW autoimmune disease.
 XX Rattus sp.
 OS
 XX W09842465-A1.
 PN
 XX 01-OCT-1998.
 PP
 XX 27-MAR-1998; 98WO-US06114.
 PP
 XX 04-SEP-1997; 97US-0057456.
 PR
 XX 27-MAR-1997; 97US-0041009.
 PR
 XX 04-JUN-1997; 97US-0869153.
 XX
 PA (1000) UNIV LOUISIANA STATE & AGRIC & MECH COLL.
 XX
 XX Elzer PL, Enright PM, Foil LD, Hansel W, Jaynes JM;
 P1 Koonce KL, McCann SM, Melrose PA, Yu WH;
 P1
 XX WPI: 1998-531711/45.
 DR
 XX
 XX New fusion of hormone and lytic peptide for selective cell killing -
 PT particularly for controlling fertility in animals, insect pests,
 PT fish etc., also for treatment of hormone-dependent cancer, viral
 PT infections and auto-immune diseases
 PT
 XX Example 1 6, Page 37; 59pp; English.
 PS
 XX This sequence represents the rat hecate peptide which is used in a method
 CC for producing long-term contraception or sterility by administering
 CC gonadotrophin-releasing hormone (GnRH), beta luteinising hormone (LH) or
 CC lampry; luteinising hormone-releasing hormone (LHRH-III) or temporarily
 CC restoring fertility in a mammal in which the pituitary gonadotropes have
 CC been selectively destroyed by administration of GnRH or LHRH-III. This
 CC sequence also has applications in methods for killing, or inhibiting
 CC growth of hormone- or ligand-dependent tumours by administering the
 CC relevant hormone- or ligand plus a lytic peptide, for killing a cell the
 CC activity of which is dependent on binding of a cell-surface receptor to
 CC a ligand by administering the ligand and lytic peptide, inhibiting
 CC reproductive capacity of an insect by administration of a lytic peptide,
 CC fusion proteins combining such proteins and lytic peptides can be fused
 CC and may also include a carrier domain, specifically vitamin B12, that
 CC facilitates intestinal uptake after oral administration. These methods
 CC have applications for increasing the growth rate of mammals, for pest
 CC control e.g. plants containing an exogenous gene encoding L-hecate. A
 CC further application is in the treatment of ovarian, breast, prostate,
 CC endometrial or testicular cancers, prolactinoma, growth hormone-,
 CC thyrotropin- or gonadotrophin-dependent adenomas, or any pituitary
 CC adenoma. In addition the proteins can be used to kill lymphocytes
 CC involved in a wide range of autoimmune diseases or to destroy
 CC virus infected cells.
 XX
 XX Sequence 23 AA;
 SQ
 Query Match 72.7%; Score 48; DB 19; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 FAFALKALKALKAL 15
 ID 1 FAFALKALKALKAL 15
 DE 1 FAFALKALKALKAL 15
 RESULT 28
 AAW49786
 ID AAW49786 standard; peptide: 23 AA.
 AC
 XX AAW49786;
 XX
 XX Amino acid sequence of a lytic peptide hecate.
 DE
 XX Contraception; sterility; amphipathic; lytic peptide; hormone domain;
 KW gonadotropin-releasing hormone; GnRH; anti-sarney; defect; defect;
 KW cow; bull; pig; horse; sheep.
 XX
 XX Synthetic.
 OS
 XX W09911282-A1.
 PN
 XX
 XX

XX 20-MAY-1998 (first entry)
 PP
 XX Synthetic peptide bp-1.
 DE
 XX lytic peptide; lysine-rich; proteolytic digestion; methylated; lysine;
 KW protection; amphipathic alpha-helix; heteroprotein; sheet; treatment;
 KW infection; viral; bacterial; yeast; fungal; protease; cancer; L-1.
 KW
 XX Synthetic.
 OS
 XX US5717064-A.
 PN
 XX 10-FEB-1998.
 PP
 XX 24-APR-1995; 95US-0427001.
 PP
 XX 24-APR-1995; 95US-0427001.
 PR
 XX 18-NOV-1993; 93US-0148889.
 PR
 XX (DEME-) DEMETER BIOTECHNOLOGIES LTD.
 PA
 XX Jaynes JM, Julian GR;
 P1
 XX WPI: 1998-168370/14.
 DR
 XX Cytolytic peptide analogues - with methylated lysine residues to
 PT increase protease resistance
 PT
 XX Example 2; Column 9; 21pp; English.
 PS
 XX Peptide AAW49796 (also known as bp-1) is used as a test substrate for a
 CC reductive alkylation reaction used in the analysis of a family of
 CC synthetic lytic peptides which are stabilised by having methylated
 CC epsilon-amino groups at their lysine residues (see AAW49796/W97965).
 CC peptides AAW49750-W49771 and AAW49781-W49785 have an amphipathic
 CC alpha-helix conformation while peptides AAW49772-W49780 have a
 CC beta-pleated sheet conformation. These peptides have enhanced resistance
 CC to proteolytic digestion by trypsin. Such peptides can be used to treat
 CC infections by lysing bacterial, yeast, fungal and protozoan cells or to
 CC treat cancers by lysing neoplastic or transformed cells. They can also be
 CC used to treat viral infections by lysing enveloped viruses and
 CC virus-infected cells.
 XX
 XX Sequence 23 AA;
 SQ
 Query Match 72.7%; Score 48; DB 19; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 FAFALKALKALKAL 15
 ID 1 FAFALKALKALKAL 15
 DE 1 FAFALKALKALKAL 15
 RESULT 29
 AAW03857
 ID AAW03857 standard; peptide: 23 AA.
 AC
 XX AAW03857;
 XX
 XX 16-JUN-1999 (first entry)
 PP
 XX Amino acid sequence of a lytic peptide hecate.
 DE
 XX Contraception; sterility; amphipathic; lytic peptide; hormone domain;
 KW gonadotropin-releasing hormone; GnRH; anti-sarney; defect; defect;
 KW cow; bull; pig; horse; sheep.
 XX
 XX Synthetic.
 OS
 XX W09911282-A1.
 PN
 XX
 XX

11-MAR 1999.
 01-SEP 1998: 98W0-US18117.
 03-SEP-1997: 97US-0057456.
 (LOUISIANA STATE & AGRIC & MECH COLL.
 Elzer PH, Enright F, Hansel W, Jaynes JM, McIlrose PA;
 WPI: 1999-204980/17.
 Production of long-term contraception or sterility in, e.g., cats
 by administering a compound comprising fusion peptide of
 gonadotropin-releasing hormone and lytic peptide domains
 Examples 1-6; Page 23; 34pp; English.
 The invention relates to method for producing long-term contraception
 or sterility in a mammal that comprises administering an amphiphilic
 lytic peptide compound. The compounds comprise: (a) a hormone domain
 selected from gonadotropin-releasing hormone (GnRH) and GnRH analogues,
 and (b) a lytic peptide domain. The compounds are relatively small and
 are not antigenic. Lysis of gonadotropes has been observed to be very
 rapid. The components: the ligand and the lytic peptide may optionally
 be administered as a fusion peptide. The methods can be applied to dogs,
 cats, cows, bulls, pigs, horses, sheep or humans, including sexually
 immature individuals.

Query Match 72.7%; Score 48; DB 20; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
 II IIII IIII I
 DI 1 FALALKALKALKAL 15

RESULT 40
 AAW84279
 11 AAW84279 standard; peptide: 23 AA.
 AAW84279;
 17 MAR 1999 (first entry)
 Lytic peptide bovine
 Hormone domain: gonadotropin releasing hormone; GnRH;
 luteinising hormone releasing hormone; LHRR-III; estrogen;
 testosterone; luteinising hormone; chorionic gonadotropin;
 follicle stimulating hormone; melanocyte-stimulating hormone; estradiol;
 dopamine; somatostatin; lytic peptide; long term contraception;
 sterility; human; fish; insect; hormone dependent tumour;
 hormone-related disease; autoimmune disorder; treatment.
 Synthetic.
 W00842794 A1.
 01 OCT 1998.
 26-MAR 1998: 98W0-US06015.
 03-SEP 1997: 97US-0057456.
 27-MAR-1997: 97US-0041009.
 04 JUN 1997: 97US-0869154.
 (DEME) DEMETER BIOTECHNOLOGIES LTD.
 (LOUISIANA STATE & AGRIC.

Enright PH, Poil LD, Hansel WB, Jaynes W, McIlrose PA;
 WPI: 1999-070064/06.
 New hormone-lytic peptide compounds used for long-term
 contraception or sterilisation, for killing disease cells, for
 infected cells or for treating tumours or other diseases.
 Example 3; Page 46; 48pp; English.
 The present sequence represents a lytic peptide (the lytic peptide) used
 in a compound in conjunction with the hormone domain of gonadotropin
 releasing hormone (GnRH), or luteinising hormone releasing hormone
 (LHRH) III, estrogen, testosterone, luteinising hormone, chorionic gonadotropin,
 gonadotropin, follicle stimulating hormone, melanocyte-stimulating hormone,
 dopamine, estradiol, dopamine, somatostatin, and melanocyte-stimulating
 hormones. The compounds can be used for specific long-term contraception
 that are driven by or are dependent on specific hormone receptors.
 They can be used for providing long-term contraception or sterility in
 humans, domesticated or wild animals, birds, reptiles, amphibians,
 bony fish, cartilaginous fish, jawless fish, invertebrates, insects,
 insects or molluscs. They can also be used for reduction of tumour
 malignant and non malignant hormone dependent tumours, for reduction of
 other hormone related diseases, for selectively killing disease cells,
 cells or for treating autoimmune disorders.

Sequence 23 AA:
 Query Match 72.7%; Score 48; DB 20; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
 II IIII IIII I
 DI 1 FALALKALKALKAL 15

RESULT 41
 APR00760
 11 ABR00760 standard; peptide: 23 AA.
 ABR00760;
 04 APR 2003 (first entry)
 Bioactive synthetic peptide; bovine AM.
 Antibacterial; tumour; cytostatic; vitiligo; cancer; reproductive system;
 genes; and immunology; human; fish; insect; melanocyte-stimulating hormone;
 bioactive.
 Synthetic.
 Key: luteinising hormone
 Modified site 23 /note "terminates with"
 W0200279408 A2.
 10 OCT 2002.
 26 MAR 2002: 2002W0-US06015.
 26 MAR 2001: 2001US 2795049.
 26 MAR 2001: 2001US 0869154.
 (DEME) DEMETER BIOTECHNOLOGIES LTD.
 Owen DR;
 WPI: 2003-231247/23.
 New isolated peptide for treating cancer, cysts or other diseases.

PT acne, inhibiting growth of microbial cells, or promoting proliferation
 PT of cells, comprises phenylalanine, leucine, alanine or lysine residues
 XX
 XX
 PS claim 7; Page 5; 13pp; English.
 XX
 CC The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicidal, cytostatic, and virostatic
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in AB000759-AB000923 represent the bioactive
 CC peptides of the invention.

XX Sequence 23 AA;

Query Match 72.7%; Score 48; DB 24; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
 II IIII IIII I
 DB 1 FALKALKALKALKL 15

RESULT 32
 ABR00945

ID ABR00945 standard; peptide: 23 AA.

AC ABR00945;

DT 03-APR-2003 (first entry)

DE Bioactive synthetic peptide Brevato AC #1010.

KW Antibacterial; fungicide; cytostatic; vulnerrary; cancer; cystic fibrosis,
 KW acne; antimicrobial; human fibroblast; human lymphocyte; wound healing,
 KW bioactive.

XX Synthetic.

PN W0200279408-A2.

XX 10-OCT-2002.

PF 28-MAR-2002; 2002WO-US09534.

XX 28-MAR-2001; 2001US-279505P.

PR 28-MAR-2001; 2001US-0820053.

PA (HELI-) HELIX BIOMEDIX INC.

XX owen DB;

XX WPI: 2003-221247/21.

PT New isolated peptide for treating cancer, cystic fibrosis, wounds or
 PT acne, inhibiting growth of microbial cells, or promoting proliferation
 PT of cells, comprises phenylalanine, leucine, alanine or lysine residues
 DT

PS claim 7; Page 5; 13pp; English.

XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicidal, cytostatic, and virostatic
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,

CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in AB000759-AB000923 represent the bioactive
 CC peptides of the invention.

XX Sequence 23 AA;

Query Match 72.7%; Score 48; DB 24; Length 23;
 Best Local Similarity 80.0%; Pred. No. 0.8;
 Matches 12; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 FAKALKALKALKAL 15
 II IIII IIII I
 DB 1 FALKALKALKALKL 15

RESULT 33

AAW07282

ID AAW07282 standard; peptide: 18 AA.

XX AAW07282;

DT 29-MAY-1997 (first entry)

XX Amphiphilic antimicrobial peptide MR 41.

XX amphiphilic antimicrobial; log kill; Staphylococcus aureus; Escherichia coli; vesicular analysis; hydrophobic lipophilic; bactericidal; hydrophobic moment; equation; antimicrobial; bactericidal; antifungal; disinfection; spore prevention; preservation.

XX Synthetic.

XX W09628468-A2.

XX 19-SEP-1996.

XX 27-FEB-1996; 96WO-EP00844.

XX 09-MAR-1995; 95GB-0004761.

AA (UNIL) UNILEVER NV.

PA (UNIL) UNILEVER PLC.

XX Bhakoo M, Patel S, Stott PJ;

XX WPI: 1996-434760/43.

PT New amphiphilic antimicrobial peptide(s) having particular
 PT combinations of hydrophilic and hydrophobic amino acid residues

XX Example -; Page 22; 39pp; English.

XX AAW07279 90 examples of amphiphilic and antimicrobial peptides for
 CC comparison to claimed amphiphilic and antimicrobial peptides (AAW 7257-90),
 CC and show that relatively minor changes in structure can have significant
 CC consequences as regards the antimicrobial activity of the molecules. The
 CC log kills were predicted against S. aureus and against E. coli and
 CC values of more than 5 and 4 respectively, correspond to greater
 CC antimicrobial activity. Effective antimicrobial peptides are
 CC discriminated from ineffective peptides by a statistical analysis in
 CC dimensions corresponding to charge, hydrophobic/lipophilic balance,
 CC hydrophobic moment and amphiphilicity of the peptides, where effective
 CC peptides fall into the region which is an envelope of the
 CC amphiphilicity dimension and clockwise of the charge dimension.
 CC Effective peptides are further discriminated from ineffective peptides by
 CC means of an equation relating certain physical properties of the peptides
 CC to their biological activity against specific microorganisms. The
 CC peptides can be used in antimicrobial, prebiotic, bactericidal or antifungal
 CC compounds. They can be used for the disinfection of surfaces, spore
 CC prevention, preservation or other hygiene processes.

XX Sequence 18 AA;

Query Match 65.28; Score 43; BH 17; Length 18;
Best Local Similarity 63.38; Pred. No. 3.6;
Matches 10; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 KALKALKALKA 14
ID 5 FALFALFALES 16

RESULT 34
AAM49118
ID AAM49118 standard; peptide: 23 AA.

XX AAM49118;
XX 20-MAY-2002 (first entry)

DE Cationic amphipathic gene transfer vector peptide; FAWA.

XX Cationic amphipathic gene transfer vector; gene therapy; non-viral;
KW nucleic acid carrier; FAWA.

XX Synthetic.

XX W0200210197-A1.

XX 07-FEB-2002.

XX 06-JUL-2001; 2001WO-0500274.

XX 20-JUL-2000; 2000ES-0001806.

XX (CNSI) CONSEJO SUPERIOR INVESTIGACIONES CIENTÍF.

PA (MEDP) MELPLANT GENETICS S.L.

PA (JESD) JESUS FORNAYA GUTIERREZ.

PA (BULEZ) BULEZ RUIZ A L.

XX Fominaya Gutierrez J., Bernad Miana A., Cassot Vega MA;

XX WP1; 2002-257378/03.

XX New cationic amphipathic peptide, useful as carrier for nucleic acid.

PI c.d. in gene therapy; binds to, and condenses, nucleic acid and

PI destabilizes cellular membranes

PS claim 5; Page 41; 54pp; Spanish.

CC The invention relates to novel cationic amphipathic peptides of the
CC general formula:
CC X1-X2-X3-X4-X5-X6-X7-X8-X9-X10-X11-X12-X13-X14-X15-X16-X17-X18-X19-X20
CC X21-X22-X23 where:
CC X1, X5, X9, X12, X19 and X23 are Arg, Lys or ornithine;
CC Val, Ileu, Ile, Trp, Tyr or Phe, with X3 preferably being an aromatic
CC amino acid, particularly Trp, the constituent amino acids of the
CC peptide may be L, or D form, and in a particular embodiment, all
CC residues are D-form. The peptides of the invention are able to
CC simultaneously bind to nucleic acid and to disrupt biological
CC membranes and may thus be used as carriers for nucleic acids,
CC particularly to prepare non-viral gene transfer vectors for gene
CC therapy or for experimental transfection. Such vectors based on
CC peptides of the invention are not inhibited by serum. Compared with
CC similar known peptides, such as KALA (AAM49119), the peptides of the
CC invention have an increased capacity for condensing genetic material,
CC have lower cytotoxicity and provide more efficient transfer of
CC genetic material. The present sequence represents a specifically
CC claimed peptide of the invention designated FAWA.

XX Sequence 23 AA;

Query Match 65.28; Score 43; BH 24; Length 23;
Best Local Similarity 64.38; Pred. No. 4.6;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 AKALKALKALKAL 15
ID 8 AKALKALKALKAL 21

RESULT 45
ABR00767
ID ABR00767 standard; peptide: 23 AA.

XX ABR00767;

XX 03-APR-2003 (first entry)

DE Bioactive synthetic peptide FLAK03 AM.

XX Anti-bacterial; fungicide; cytostatic; without D-tryptophan; without D-phenylalanine;
KW anti-infective; human fibrinolytic; human angiogenic; human bone marrow
XX bioactive.

XX Synthetic.

XX Key location/qualifiers

XX Modified site 23

XX Note: "C" terminal amide "

XX W0200279408 A2.

XX 10-OCT-2002.

XX 28-MAR-2002; 2002WO-US09654.

XX 28-MAR-2001; 2001US-279408A.

XX 28-MAR-2001; 2001US-062006A.

XX ABEL / HELL / MED-IX INC.

XX Owen DR;

XX W01; 2003-223247/21.

XX New isolated peptide for treating cancer, especially for inhibiting tumor growth,
PI inhibiting growth of microbial cells, or promoting proliferation of cells,
PI comprises phenylalanine, leucine, isoleucine or valine residues.
XX Example 25; Page 5; 13pp; English.

CC The invention relates to a novel isolated peptide which is a linear chain of
CC acids in length, and comprises phenylalanine, leucine, isoleucine or valine
CC residues, or contains at least one of the residues. The peptide of the
CC invention have anti-bacterial, fungicide, cytostatic, and cancer
CC activity. The peptides are useful in treatment of cancer, especially in
CC cancer, inhibiting the growth of microbial cells, promoting proliferation of
CC stimulation and/or proliferation of human cells, and inhibiting tumor
CC promoting wound healing, and in enhancing the immune system of a patient.
CC present. The sequences shown in ABR00767 and W0200279408 A2 are the
CC peptides of the invention.

XX Sequence 23 AA;

Query Match 65.28; Score 43; BH 24; Length 23;
Best Local Similarity 74.89; Pred. No. 3.0;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 PARAKALKALKAL 14
ID 1 PARAKALKALKAL 15

RESULT 46
ABR00777
ID ABR00777 standard; peptide: 23 AA.

XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicide, cytostatic, and vulnerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC stimulating wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in ABR00759 ABR00923 represent the bioactive
 CC peptides of the invention.

XX Sequence 23 AA:

Query Match 63.6% Score 42; DB 24; Length 23;
 Best Local Similarity 78.6%; Pred. No. 6.5;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AAALFALEAKAL 15
 1 | | | | | | | | |
 DB 2 AAALFALEAKAL 15

RESULT 41

ABR00916
 ID ABR00916 standard; peptide: 23 AA.

AC ABR00916;

XX 03-APR-2003 (first entry)

XX Bioactive synthetic peptide HECATE AMV.

XX Antibacterial; fungicide; cytostatic; vulnerary; cancer; cystic fibrosis;
 KW acne; antimicrobial; human fibroblast; human lymphocyte; wound healing;
 KW bioactive.

XX Synthetic.

XX Key Location/Qualifiers

XX Modified-site 24 /note: "C-terminal amide"

XX W020279438 A2.

XX 10-OCT-2002.

XX 28-MAR-2002; 2002WO-US09534.

XX 28-MAR-2001; 2001US-274505P.

XX 28-MAR-2001; 2001US-0820053.

XX (HELI-) HELIX BIOMEDIX INC.

XX Owen DR;

XX WPI; 2003 221217/21.

XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
 CC acne, inhibiting growth of microbial cells, or promoting proliferation
 CC of cells, comprises phenylalanine, leucine, alanine or lysine residues

XX Example 2; Page 9; 13pp; English.

XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicide, cytostatic, and vulnerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic

CC agent. The sequences shown in ABR00759 ABR00923 represent the bioactive
 CC peptides of the invention.

XX Sequence 23 AA:

Query Match 63.6% Score 42; DB 24; Length 23;
 Best Local Similarity 78.6%; Pred. No. 6.5;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AAALFALEAKAL 15
 1 | | | | | | | | |
 DB 2 AAALFALEAKAL 15

RESULT 42

ABR00917
 ID ABR00917 standard; peptide: 23 AA.

AC ABR00917;

XX 03-APR-2003 (first entry)

XX Bioactive synthetic peptide HECATE ACV.

XX Antibacterial; fungicide; cytostatic; vulnerary; cancer; cystic fibrosis;
 KW acne; antimicrobial; human fibroblast; human lymphocyte; wound healing;
 KW bioactive.

XX Synthetic.

XX W0200279408 A2.

XX 10-OCT-2002.

XX 28-MAR-2002; 2002WO-US09534.

XX 28-MAR-2001; 2001US-274505P.

XX 28-MAR-2001; 2001US-0820053.

XX (HELI-) HELIX BIOMEDIX INC.

XX Owen DR;

XX WPI; 2003 221217/21.

XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
 CC acne, inhibiting growth of microbial cells, or promoting proliferation
 CC of cells, comprises phenylalanine, leucine, alanine or lysine residues

XX Claim 7; Page 9; 13pp; English.

XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicide, cytostatic, and vulnerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in ABR00759 ABR00923 represent the bioactive
 CC peptides of the invention.

XX Sequence 23 AA:

Query Match 63.6% Score 42; DB 24; Length 23;
 Best Local Similarity 78.6%; Pred. No. 6.5;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 AAALFALEAKAL 15
 1 | | | | | | | | |
 DB 2 AAALFALEAKAL 15

```

RESULT 43
ABR00760 standard: peptide: 15 AA.
XX ABR00760;
XX
XX 03-APR-2003 (first entry)
XX
XX Bioactive synthetic peptide FLAK62 AM.
XX
XX Antibacterial: fungicide; cystostatic; cytotoxic; cancer; cystic fibrosis;
XX active antimicrobial; human fibroblast; human lymphocyte; wound healing;
XX Bioactive.
XX
XX Synthetic.
XX
XX Key location/Qualifiers
XX Modified-site 15
XX /note: "C-terminal amide"
XX
XX W220279436 A2.
XX 10-OCT-2002.
XX
XX 28-MAR-2002: 2002WO-US09544.
XX
XX 28-MAR-2001: 2001US-279505P.
XX 28-MAR-2001: 2001US-082005A.
XX
XX (HELI) HELIX BIOMEDIX INC.
XX
XX Owen DR.
XX
XX WPI: 2003-221247/21.
XX
XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
XX acne, inhibiting growth of microbial cells, or promoting proliferation
XX of cells, comprises phenylalanine, leucine, alanine or lysine residues
XX
XX Example 2: Page 6: 134pp: English.
XX
XX The invention relates to a novel isolated peptide which is 5-23 amino
XX acids in length, and comprising phenylalanine, leucine, alanine or lysine
XX residues, or contains at least 50 % of the residues. The peptides of the
XX invention have antibacterial, fungicide, cytostatic, and cytotoxic
XX activity. The peptides are useful in treating cancer, cystic fibrosis or
XX acne, inhibiting the growth of microbial cells, promoting the
XX stimulation and/or proliferation of human fibroblasts and lymphocytes,
XX promoting wound healing, and in enhancing the activity of a therapeutic
XX agent. The sequences shown in ABR00760 ABR00764 represent the bioactive
XX peptides of the invention.
XX
XX Sequence 15 AA:
XX
XX Query Match 62.1% Score 41: DB 24: Length 15;
XX Best Local Similarity 74.4% Pred. No. 6;
XX Matches 11: Conservative 0: Mismatches 4: Indels 0: Gaps 0;
XX
XX 1 FAKAKKALKKALK 15
XX 1 FAKAKKALKKALK 15
XX
XX RESULT 44
ABR00764 standard: peptide: 16 AA.
XX ABR00764;
XX
XX 03-APR-2003 (first entry)
XX
XX Bioactive synthetic peptide FLAK62 AMV.
XX
XX Antibacterial: fungicide; cystostatic; cytotoxic; cancer; cystic fibrosis;
XX active antimicrobial; human fibroblast; human lymphocyte; wound healing;
XX Bioactive.
XX
XX Synthetic.
XX
XX Key location/Qualifiers
XX Modified-site 16
XX /note: "C-terminal amide"
XX
XX W220279436 A2.
XX 10-OCT-2002.
XX
XX 28-MAR-2002: 2002WO-US09544.
XX
XX 28-MAR-2001: 2001US-279505P.
XX 28-MAR-2001: 2001US-082005A.
XX
XX (HELI) HELIX BIOMEDIX INC.
XX
XX Owen DR.
XX
XX WPI: 2003-221247/21.
XX
XX New isolated peptide for treating cancer, cystic fibrosis, wounds or
XX acne, inhibiting growth of microbial cells, or promoting proliferation
XX of cells, comprises phenylalanine, leucine, alanine or lysine residues
XX
XX Claim 2: Page 6: 134pp: English.
XX
XX the invention relates to a novel isolated peptide which is 5-23 amino
XX acids in length, and comprising phenylalanine, leucine, alanine or lysine
XX residues, or contains at least 50 % of the residues. The peptides of the
XX invention have antibacterial, fungicide, cytostatic, and cytotoxic
XX activity. The peptides are useful in treating cancer, cystic fibrosis or
XX acne, inhibiting the growth of microbial cells, promoting the
XX stimulation and/or proliferation of human fibroblasts and lymphocytes,
XX promoting wound healing, and in enhancing the activity of a therapeutic
XX agent. The sequences shown in ABR00760 ABR00764 represent the bioactive
XX peptides of the invention.
XX
XX Sequence 16 AA:
XX
XX Query Match 62.1% Score 41: DB 24: Length 16;
XX Best Local Similarity 76.9% Pred. No. 6;4;
XX Matches 10: Conservative 0: Mismatches 4: Indels 0: Gaps 0;
XX
XX 1 FAKAKKALKKALK 16
XX 1 FAKAKKALKKALK 16
XX
XX RESULT 45
ABR00769 standard: peptide: 24 AA.
XX ABR00769;
XX
XX 03-APR-2003 (first entry)
XX
XX Bioactive synthetic peptide FLAK62 AMV.
XX
XX Antibacterial: fungicide; cystostatic; cytotoxic; cancer; cystic fibrosis;
XX active antimicrobial; human fibroblast; human lymphocyte; wound healing;
XX Bioactive.
XX
XX Synthetic.
XX
XX Key location/Qualifiers

```

FT Modified-site 24 /note- "C-terminal amide"

XX WO200279408-A2.

XX 10-OCT-2002.

XX 28-MAR-2002; 2002WO-0509534.

XX 28-MAR-2001; 2001US-279505P.

XX 28-MAR-2001; 2001US-0820053.

XX (HELI) HELIX BIOMEDIX INC.

XX OWON DR;

XX WP; 2002 22124721.

PT Now isolated peptide for treating cancer, cystic fibrosis, wounds or
 PT acne, inhibiting growth of microbial cells, or promoting proliferation
 PT of cells, comprises phenylalanine, leucine, alanine or lysine residues
 PT

XX Example 2: Page 9; 13pp; English.

XX The invention relates to a novel isolated peptide which is 5-23 amino
 CC acids in length, and comprising phenylalanine, leucine, alanine or lysine
 CC residues, or contains at least 50 % of the residues. The peptides of the
 CC invention have antibacterial, fungicide, cytostatic, and vulcerary
 CC activity. The peptides are useful in treating cancer, cystic fibrosis or
 CC acne, inhibiting the growth of microbial cells, promoting the
 CC stimulation and/or proliferation of human fibroblasts and lymphocytes,
 CC promoting wound healing, and in enhancing the activity of a therapeutic
 CC agent. The sequences shown in AB00759-AB00923 represent the bioactive
 CC peptides of the invention.

XX Sequence 24 AA:

Query Match 62.1%; Score 41; DB 24; Length 22;
 Best Local Similarity 76.9%; Pred. No. 9.3;
 Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 KALKALKALKAL 15

DB 6 KALKALKALKAL 18

|||||

Search completed: August 21, 2003, 08:19:04

Job Time : 84 secs



GenCore version 5.1.6
Copyright (c) 1993-2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: August 21, 2003, 08:08:00 ; Search time 94 Seconds
(without alignments)
41.179 Million cell updates/sec

Title: us-09-820-053a-43
Perfect score: 66
Sequence: 1 FAKALKALKAL 15

Scoring table: BLOSUM62
Gapop 10.0 ; Gapov 0.5

Searched: 840525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 10083

Minimum DB seq length: 0
Maximum DB seq length: 25

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMHU_21:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_orqanelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_virus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	29	43.9	20	2 Q56130	Q56130 salmonella
2	26	39.4	21	2 Q46010	Q46010 clostridium
3	26	39.4	21	2 Q46011	Q46011 clostridium
4	26	39.4	25	15 Q99828	Q99828 human lamal
5	26	39.4	25	15 Q99826	Q99826 human lamal
6	26	39.4	25	15 Q99817	Q99817 human lamal
7	25	37.9	18	1 Q90827	Q90827 methanococcus
8	24	36.4	11	2 P77404	P77404 escherichia
9	24	36.4	11	2 P95518	P95518 pasteurella
10	24	36.4	19	2 Q99510	Q99510 enterobacter
11	24	36.4	20	13 P82876	P82876 rana clamit
12	24	36.4	24	5 Q9N685	Q9N685 strongylova
13	23	34.8	15	13 P83333	P83333 oncorhynch
14	23	34.8	17	8 Q93468	Q93468 brassica ju
15	23	34.8	19	4 Q99711	Q99711 homo sapien
16	23	34.8	23	12 Q97050	Q97050 pseudotritic

ALIGNMENTS

RESULT 1

Q56130 ID Q56130 PRELIMINARY: PRI: 20 AA.
AC Q56130:
DI 01-NOV-1996 (JEMBLrel_01, Created)
DI 01-NOV-1996 (JEMBLrel_01, Last sequence update)
DI 01-DEC-2001 (JEMBLrel_19, Last annotation update)
DE typhimurium rpoB RNA polymerase beta subunit (Fragment).
OS Salmonella typhimurium.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID:602;
RN (1)
RP SEQUENCE FROM N.A.
EA Svetlana E.P., Litsyn N.A., Gurev S.G., Muzlytskaya G.S.;
RT "Nucleotide sequence of the rpoB gene of Salmonella typhimurium.
RT Coding for the beta subunit of RNA polymerase.";
RL Dck1, Biochem. 287:232-246(1986).
RN (2)
RP SEQUENCE FROM N.A.
KA Svetlana E.P., Litsyn N.A., Gurev S.G., Muzlytskaya G.S.;
RT "Nucleotide sequence of the rpoB gene of Salmonella typhimurium.
RT for the beta subunit of RNA polymerase.";
RL Dck1, Biochem. 287:62-65(1986).
RN (3)
RP SEQUENCE FROM N.A.
KA Svetlana E.P., Litsyn N.A., Muzlytskaya G.S.;
RT "Sequences coding for RNA polymerase beta subunit (in bacteria).";
RL Dck1, Biochem. 177:563-569(1986).
RN (4)
RP NON-REF
SQ Q99826: 20 20 20
P82876: 20 AA, 2376 MW, 6246.443 aa, 677A, 669647

Query Match 43.9% Score 29.0
Best local similarity 70.9% Pred. No. 5, seq. 2;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0

us-09-820-053a-43


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ID Q00017 PRELIMINARY: PRI: 25 AA.
AC Q00017
DT 01-MAR-2001 (TEMBLrel. 16, Created)
DI 01-MAY-2001 (TEMBLrel. 15, Last sequence update)
DE 01-OCT-2002 (TEMBLrel. 22, Last annotation update)
RE Rev protein. (Fragment).
GN REV.
OS Human immunodeficiency virus 1
OC Retrovira; Retrovira; Retroviridae; Lentivirus.
OX NCBI_TaxID 11676;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 21002574; PubMed: 11118069;
RA Mollmura F., Fuchida S., Fato R., Mochizuki K., Oishi H.K., Tani K.K.,
RT Thwe M., Oo K.Y., Lwin S., Kyaw O., Zaw M., Nappi Y., Takebe Y.;
RT "Emergence of new forms of human immunodeficiency virus type 1
RT intersubtype recombinants in central Myanmar."
RL AIDS Res. Hum. Retroviruses 16:1831-1843(2000).
DR EMBL: AB043904; BAB19238.1;
DR InterPro: IPR000625; REV_protein.
DR Pfam: PF00424; REV: 1.
FI NCBI_TaxID: 25.
SI SEQUENCE: 25 AA: 2735 MW: 45933.93AA:412248 CR:64.

Query Match: 89.4%; Score 26; DB 15; Length 25;
Best Local Similarity: 50.0%; Prod. No. 2.1e+04;
Matches: 5; Conservative: 4; Mismatches: 1; Indels: 0; Gaps: 0;

QY 6 KALLKALKAL 15
DB 10 EALLKAVRI 19

RESULT 7
Q00017 PRELIMINARY: PRI: 18 AA.
ID Q00017
AC Q00017
DT 01-MAY-2000 (TEMBLrel. 14, Created)
DI 01-MAY-2000 (TEMBLrel. 14, Last sequence update)
DE 01-JUN-2000 (TEMBLrel. 14, Last annotation update)
DE METHYLOHALAMIN: coenzyme M methyltransferase isoenzyme II (Fragment).
OS Methanosarcina barkeri.
OC Archaea; Euryarchaeota; Methanococci; Methanosarcinales;
OC Methanosarcinaceae; Methanosarcina.
OX NCBI_TaxID: 2208;
RN [1]
RP SEQUENCE.
RX MEDLINE: 94356596; PubMed: 8252643;
RA Velissov A., Gartner P., Harris D., Linder D., Thauer R.K.;
RT "Function of methylcobalamin: coenzyme M methyltransferase isoenzyme
RT II in Methanosarcina barkeri."
RL Arch. Microbiol. 159:530-536(1994).
SI SEQUENCE: 18 AA: 1916 MW: 180442564322875 CR:64;

Query Match: 87.0%; Score 25; DB 1; Length 18;
Best Local Similarity: 66.7%; Prod. No. 2.2e+03;
Matches: 6; Conservative: 1; Mismatches: 2; Indels: 0; Gaps: 0;

QY 5 LKALLKALK 13
DB 5 LKTLAALE 14

RESULT 8
P77404 PRELIMINARY: PRI: 11 AA.
ID P77404
AC P77404
DT 01-FEB-1997 (TEMBLrel. 02, Created)
DI 01-FEB-1997 (TEMBLrel. 02, Last sequence update)
DI 01-NOV-1998 (TEMBLrel. 08, Last annotation update)
DE DNA sequence downstream of the EOPKRI HSD locus (Fragment).
GN HSDR.

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OS Escherichia coli.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriaceae;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID 562;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 97296151; PubMed: 9157244;
RA Eyo J.L., C. Johnson B., Sridharan P., Kulkarni R., Bhat K.V.,
RT "The type 1 and 2 of the enterobacterial enterohemorrhagic EHEC with
RT high homology to the phage phi genome: implications for the evolution
RT and spread of DNA restriction systems."
RL Mol. Microbiol. 24:729-736(1997).
DR EMBL: X98145; CAA66840.1;
DR EMBL: X98144; CAA66840.1;
FI NCBI_TaxID: 1.
SI SEQUENCE: 11 AA: 1259 MW: 71469925432154 CR:54;

Query Match: 66.4%; Score 24; DB 2; Length 11;
Best Local Similarity: 71.4%; Prod. No. 1.7e+03;
Matches: 5; Conservative: 2; Mismatches: 3; Indels: 0; Gaps: 0;

QY 8 LKALKAKA 14
DB 5 LKLSLKA 11

RESULT 9
P95518 PRELIMINARY: PRI: 11 AA.
ID P95518
AC P95518
DT 01-MAY-1997 (TEMBLrel. 03, Created)
DI 01-MAY-1997 (TEMBLrel. 03, Last sequence update)
DI 01-DEC-2001 (TEMBLrel. 19, Last annotation update)
DE Ribosomal protein RpsA (Fragment).
GN RPSA.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellales; Mannheimia.
OX NCBI_TaxID 75985;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN DRI101;
RX MEDLINE: 9716447; PubMed: 9011048;
RA Hindinger S.K., Garza G., Brown R.J., Kady S.;
RT "Isolation and characterization of the intracellular host factor species
RT of Pasteurella haemolytica."
RL FEMS Microbiol. Lett. 146:183-188(1997).
DR EMBL: 056149; AAC44845.1;
FI NCBI_TaxID: 1.
SI SEQUENCE: 11 AA: 1168 MW: 7446800000000000 CR:54;

Query Match: 66.4%; Score 24; DB 2; Length 11;
Best Local Similarity: 71.4%; Prod. No. 1.7e+03;
Matches: 5; Conservative: 1; Mismatches: 3; Indels: 0; Gaps: 0;

QY 7 FAKAKAKA 7
DB 2 FAKAKAKA 8

RESULT 10
Q98519 PRELIMINARY: PRI: 19 AA.
ID Q98519
AC Q98519
DT 01-MAY-2000 (TEMBLrel. 14, Created)
DI 01-MAY-2000 (TEMBLrel. 14, Last sequence update)
DI 01-JUN-2000 (TEMBLrel. 14, Last annotation update)
DE L-2,4-diaminobutyrate decarboxylase.
OS Acetobacter calcoaceticus.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Escobacteriaceae;
OC Moraxellaceae; Acetobacter.
OX NCBI_TaxID: 471;
RN [1]

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OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eucosids 11; Brassicales; Brassicaceae; Brassica;
OX NCBI_TaxID=3707;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN cv. Zhebond no. 1;
RC "Chen X.J.; Zhang Y.Z.; Chen Z.J.; Jin Y.;
RT "A specific mtDNA fragment related to cytoplasmic male sterility in
RT tuber mustard.";
RL Submitted (Aug 2003) to the EMBL Genbank, DDBJ databases.
DR EMBL: AF298549; AAG49440.1;
KW Hypothetical protein; Mitochondrion.
SQ SEQUENCE 17 AA, 2019 MW, 55670CDRFE69EEZ6 CR6b4;
Query Match 34.8%; Score 23; DB 8; Length 17;
Best Local Similarity 55.6%; Pred. No. 4.3e+03;
Matches 5; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
QY 4 KALKALKL 11
DB [1]
DB 8 KQKMLVKS 16
[1]
RESULT 15
Q99711 PRELIMINARY; PRT; 19 AA.
ID Q99711;
AC Q99711;
DT 01-MAY-1997 (FEBRELEL_03, Created)
DI 01-MAY-1997 (FEBRELEL_03, last sequence update)
DE 01-DEC-2001 (FEBRELEL_19, last annotation update)
DE Gamma-glutamylcysteine synthetase light subunit (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE 98278973; PubMed=9614065;
RA Molnava H.R.; Mulcahy R.L.;
RT "An electrophile responsive element (EpRE) regulates beta-
RT naphthoflavone induction of the human gamma-glutamylcysteine
RT synthetase regulatory subunit gene. Constitutive expression is
RT mediated by an adjacent ap-1 site.";
RL J. Biol. Chem. 273:14683-14689(1998).
DR EMBL: U72210; AAC24417.1;
FT NON_CODING 19
SQ SEQUENCE 19 AA, 1993 MW, 5E379EA26886D15C CR6b4;
Query Match 34.8%; Score 23; DB 4; Length 19;
Best Local Similarity 62.5%; Pred. No. 4.8e+03;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 2 AKALKALL 9
DB [1]
DB 5 SRAAKALL 12
[1]
RESULT 16
Q87078 PRELIMINARY; PRT; 23 AA.
ID Q87078;
AC Q87078;
DT 01-NOV-1996 (FEBRELEL_01, Created)
DI 01-NOV-1996 (FEBRELEL_01, last sequence update)
DI 01-NOV-1998 (FEBRELEL_08, last annotation update)
DE Glycoprotein c precursor (Fragment).
OS Pseudorabies virus.
OC Viruses; dsRNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OX NCBI_TaxID=10345;
RN [1]
RP SEQUENCE FROM N.A.

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RC STRAIN Becker;
RC MEDLINE 96070924; PubMed 7499261;
RA Ryan P.; Edwards C.O.;
RT "Systematic introduction of proline in a eukaryotic signal sequence
RT suggests asymmetry within the hydrophobic core.";
RL J. Biol. Chem. 270:27876-27879(1995).
DR EMBL: U29121; AAC54531.1;
KW Signal.
FT SIGNAL 1 22 POTENTIAL.
FT NON_CODING 23
SQ SEQUENCE 23 AA, 2229 MW, 67112b1121437473 CR6b4;
Query Match 34.8%; Score 23; DB 12; Length 23;
Best Local Similarity 62.5%; Pred. No. 5.8e+03;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 2 AKALKALL 9
DB [1]
DB 5 ARAMPALL 12
[1]
RESULT 17
Q99BR3 PRELIMINARY; PRT; 25 AA.
ID Q99BR3;
AC Q99BR3;
DT 01-JUN-2001 (FEBRELEL_17, Created)
DI 01-JUN-2001 (FEBRELEL_17, last sequence update)
DI 01-OCT-2002 (FEBRELEL_22, last annotation update)
DE Rev protein (Fragment).
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retroviruses; Retroviridae; Lentivirus.
OX NCBI_TaxID=11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN TV005-R1;
RC MEDLINE 21322026; PubMed=11429118;
RA Scriba T.J.; Friedman R.K.; Zeller M.; Enderbrecht S.;
RA van Kesteren E.J.;
RT "Characterization and phylogenetic analysis of South African HIV-1
RT subtype 3 accessory genes.";
RL AIDS Res. Hum. Retroviruses 17:776-781(2001).
DR EMBL: AF425747; AAK09125.1;
DR Infoprot: IPR000625; REV_protein.
FT PROTEIN 25
FT NON_CODING 25
SQ SEQUENCE 25 AA, 2723 MW, 12933F144E1225A CR6b4;
Query Match 34.8%; Score 23; DB 15; Length 25;
Best Local Similarity 40.0%; Pred. No. 6.9e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;
QY 6 KALKALKALL 16
DB [1]
DB 10 EALLSAVRRI 19
[1]
RESULT 18
Q99BS6 PRELIMINARY; PRT; 25 AA.
ID Q99BS6;
AC Q99BS6;
DT 01-JUN-2001 (FEBRELEL_17, Created)
DI 01-JUN-2001 (FEBRELEL_17, last sequence update)
DI 01-OCT-2002 (FEBRELEL_22, last annotation update)
DE Rev protein (Fragment).
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retroviruses; Retroviridae; Lentivirus.
OX NCBI_TaxID=11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN TV004_45;
RC MEDLINE 21322026; PubMed 11429118;

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[illegible]

Matches 6: Conservative 0: Mismatches 3: Indels 0: Gaps 0:

QY 2 AKALKALK 10
11111111
DB 4 AKALKALK 12

RESULT 23

Q14001
ID Q14001 PRELIMINARY: PRI: 17 AA.
AC Q14001
DT 01 NOV 1996 (TREMBLER, 01, Created)
DT 01 NOV 1996 (TREMBLER, 01, Last sequence update)
DT 01 DEC 2001 (TREMBLER, 19, Last annotation update)
DE Cyclic nucleotide phosphodiesterase (Fragment).
GN CIPHER.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
OX NCBI TaxID 9606;
RN 11
RP SEQUENCE FROM N.A.
RX MEDLINE 97074687; PubMed 8921498;
RA Leichter R.W., Winterpacht A., Seipel R., Zabel R.D.;
RF "Molecular cloning and chromosomal assignment of the human homologue
R1 of the rat cAMP-inhibited phosphodiesterase 1 (pDEA) A gene involved
R1 in fat metabolism located at 11p15.1.";
RL Genomics 87:211-218(1996).
DR EMBL: X95522; CAA64776.1;
FI NON_TER 17
SQ SEQUENCE 17 AA: 2057 MW: 65011734FAE11540 GKCG4.

Query Match 33.48; Score 22; DB 2; Length 17;
Best Local Similarity 57.18; Pred. No. 6, to 03;
Matches 4: Conservative 3: Mismatches 0: Indels 0: Gaps 0:

QY 2 AKALKALK 9
11111111
DB 8 AKAMPSL 14

RESULT 24

Q26342
ID Q26342 PRELIMINARY: PRI: 18 AA.
AC Q26342
DT 01 MAY 1999 (TREMBLER, 10, Created)
DT 01 MAY 1999 (TREMBLER, 10, Last sequence update)
DE Protein transport protein (Fragment).
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiae; Chlamydiales; Chlamydiaceae; Chlamydia.
OX NCBI TaxID 81;
RN 11
RP SEQUENCE FROM N.A.
RX STRAIN L2 4349;
RA Wang L., Steenborg S.D., Zheng Y., Larson S.H.;
RF "Gene identification of chlamydia trachomatis by random DNA
R1 sequencing.";
RL Submitted (Aug-1998) to the EMBL/Genbank/DBJ databases.
DR EMBL: AF067425; AA004100.1;
FI NON_TER 18
SQ SEQUENCE 18 AA: 2236 MW: 5503A82ED42C0201 GKCG4.

Query Match 33.48; Score 22; DB 2; Length 18;
Best Local Similarity 83.88; Pred. No. 6, to 04;
Matches 5: Conservative 0: Mismatches 1: Indels 0: Gaps 0:

QY 1 FAKALK 6
111111
DB 10 FYEALK 15

RESULT 25

Q984P1
ID Q984P1 PRELIMINARY: PRI: 19 AA.
AC Q984P1
DT 01 MAY 2000 (TREMBLER, 13, Created)
DT 01 MAY 2000 (TREMBLER, 13, Last sequence update)
DT 01 JUN 2000 (TREMBLER, 14, Last annotation update)
DE Ribosomal protein S21 (Fragment).
OS Brucella abortus (Brucellaceae; Alphaproteobacteria; Gammaproteobacteria).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Gammaproteobacteria;
OC Caulobacteraceae; Brucellaceae.
OX NCBI TaxID 41276;
RN 11
RP SEQUENCE.
RX MEDLINE 95244409; PubMed 7727274;
RA Cell K.;
RF "Comparative ribosomal protein sequence analyses for 123 ribosomal
R1 defined atoms, Brucellaceae, and 123 other species.";
RL Int. J. Syst. Bacteriol. 45:208-215(1995).
SQ SEQUENCE 19 AA: 2071 MW: 226647514P41E5K 36042

Query Match 33.48; Score 22; DB 2; Length 19;
Best Local Similarity 83.88; Pred. No. 6, to 04;
Matches 5: Conservative 1: Mismatches 0: Indels 0: Gaps 0:

QY 1 FAKALK 8
111111
DB 12 GAKALK 17

RESULT 26

Q9SB18
ID Q9SB18 PRELIMINARY: PRI: 19 AA.
AC Q9SB18
DT 01 MAY 2000 (TREMBLER, 13, Created)
DT 01 MAY 2000 (TREMBLER, 13, Last sequence update)
DT 01 SEP 2001 (TREMBLER, 19, Last annotation update)
DE Potato patatin (Fragment).
OS Solanum tuberosum (Potato).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Charophyta;
OC Spermatophyta; Magnoliophyta; Eudicotyledons; Eucomiales; Solanales;
OC Asteridae; Lamiales; Solanales; Solanaceae; Solanum.
OX NCBI TaxID 4113;
RN 11
RP SEQUENCE FROM N.A.
RX Isoli D., Ooms G.;
RF "The 5' flanking DNA of a patatin gene directs a tissue-specific
R1 expression of a chimeric gene in potato.";
RL Plant Mol. Biol. 9:445-475(1987).
LR EMBL: M17640; AAA34818.1;
FI NON_TER 19
SQ SEQUENCE 19 AA: 2109 MW: 12097A90007D6243B544

Query Match 33.48; Score 22; DB 2; Length 19;
Best Local Similarity 50.00; Pred. No. 6, to 05;
Matches 5: Conservative 2: Mismatches 4: Indels 0: Gaps 0:

QY 1 FAKALK KALKALK 14
111111111111
DB 1 FAKALKSPEKLEAKNEA 19

RESULT 27

Q90K05
ID Q90K05 PRELIMINARY: PRI: 19 AA.
AC Q90K05
DT 01 MAY 2000 (TREMBLER, 13, Created)
DT 01 MAY 2000 (TREMBLER, 13, Last sequence update)
DT 01 MAY 2000 (TREMBLER, 13, Last annotation update)
DE Voltage-gated sodium channel NaV (Fragment).
GN CERNIA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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OX Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.
RN [11]
RP SEQUENCE FROM N.A.
RX MEDLINE 99475424; PubMed 10444332;
RA DDB Bait S.D., Tyrrell L., Escaya A., Wood P.M., Meisler M.H.,
KA Waxman S.G.
R1 "Coding sequence, genomic organization, and conserved chromosomal
R2 localization of the mouse gene Scn1a encoding the sodium channel
R3 Nav1.1."
R4 Genomics 59:409-418(1999).
DR EMBL AF126749; AAS54407.1;
KW Ionic channel.
FT NON_TER 1 19
FT NON_TER 19 19
SQ SEQUENCE 19 AA: 2174 MW: 260342339414374 (RC64);
Query Match 33.38; Score 22; DB 4; Length 19;
Best Local Similarity 50.0%; Pred. No. 75003;
Matches 4; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 8 LKALKAL 15
DB 8 VERALKAL 15
[11]
RESULT 29
Q98R76 PRELIMINARY: PRI: 20 AA.
ID Q98R76
AC Q98R76
DI 01-JUN-2001 (FIREBASEL 17, Created)
DI 01-JUN-2001 (FIREBASEL 17, Last sequence update)
DI 01-JUN-2001 (FIREBASEL 17, Last annotation update)
DE Peroxisomal membrane protein 1 (Fragment).
CN ABC13.
OS Sus scrofa (pig).
OC Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:
OC Mammalia: Eutheria: Catarrhini: Suidae: Sus.
OX NCBI TaxID 9623;
RN [11]
RP SEQUENCE FROM N.A.
RA STRAIN Protein.
KA Strull A., Kubiakova S., Poolman L.J., Reiner G., Musilova P.,
KA Van Poucke M., Robes J., Goldmann H.
R1 "FISH, RFL and linkage assignment of the porcine Abp1 (PMP1) gene to
R2 the distal end of chromosome 4q."
R3 Submitted (2001-09-11) to the EMBL/GenBank/Tran database.
DR EMBL AL009827; CAC32854.1;
FT NON_TER 1 20
FT NON_TER 20 20
SQ SEQUENCE 20 AA: 2215 MW: 257197520805088 (RC64);
Query Match 33.38; Score 22; DB 6; Length 20;
Best Local Similarity 33.38; Pred. No. 74000;
Matches 5; Conservative 4; Mismatches 7; Indels 0; Gaps 0;
QY 1 PAKALKALKAL 16
DB 1 PAKALKALKAL 16
[11]
RESULT 29
Q98R77 PRELIMINARY: PRI: 20 AA.
ID Q98R77
AC Q98R77
DI 01-MAR-2001 (FIREBASEL 16, Created)
DI 01-MAR-2001 (FIREBASEL 16, Last sequence update)
DI 01-MAR-2001 (FIREBASEL 16, Last annotation update)
DE Rana clamata (green frog).
OS Rana clamata (green frog).
OC Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:
OC Amphibia: Batrachia: Anura: Neobatrachia: Ranoidae: Rana.
OX NCBI TaxID 145282;

```

```

RN [11]
RP SEQUENCE.
RX MEDLINE 2026065; PubMed 10622101;
RA Balverson L., Basky Y.L., Kuo P.T., Campbell M.
R1 "Partial cloning and characterization of an intracellular loop of a
R2 skin of the North American green tree frog (Rana clamata)."
R3 peptides 21469.476(2000).
R4 "THE FIRST ANTIHERPESIN ACTIVITY AGAINST HSV-1 IS LOCATED IN THE
R5 C-TERMINUS AND GRAM NEGATIVE BACTERIUM FUS11. HA ACTIVELY AGAINST
R6 C. ALBICANS (BY SIMILARITY).
R7 SUBCELLULAR LOCATION: SECRETED.
R8 MASS SPECTROMETRY: MW 2109.0; MW 2109.0; MW 2109.0; MW 2109.0; MW
R9 SIMILARITY: BELONGS TO THE RELVING/RELVING/RELVING/RELVING/RELVING
R0 FAMILY.
KW Anti-herpesin: Fungicide.
FT DISULFID 14 20
SQ SEQUENCE 20 AA: 2112 MW: 2021062706027176(94);
Query Match 33.38; Score 22; DB 14; Length 20;
Best Local Similarity 45.58; Pred. No. 74000;
Matches 5; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
QY 5 LKALKALKAL 16
DB 2 LKALKALKAL 16
[11]
RESULT 40
Q98R50 PRELIMINARY: PRI: 20 AA.
ID Q98R50
AC Q98R50
DI 01-OCT-2002 (FIREBASEL 22, Created)
DI 01-OCT-2002 (FIREBASEL 22, Last sequence update)
DI 01-OCT-2002 (FIREBASEL 22, Last annotation update)
DE Hypothetical protein (Fragment).
OS Homo sapiens (human).
OC Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:
OC Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.
OX NCBI TaxID 9606;
RN [11]
RP SEQUENCE FROM N.A.
RA STRAIN Lung.
KA Strain Lung.
R1 "Submitted (2001-09-11) to the EMBL/GenBank/Tran database.
R2 EMBL AL009827; CAC32854.1;
R3 Hypothetical protein.
FT NON_TER 1 1
FT NON_TER 1 1
SQ SEQUENCE 21 AA: 2203 MW: 22021062706027176(94);
Query Match 33.38; Score 22; DB 14; Length 21;
Best Local Similarity 45.58; Pred. No. 74000;
Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
QY 2 AKALKALKAL 14
DB 2 AKALKALKAL 14
[11]
RESULT 41
Q98R52 PRELIMINARY: PRI: 20 AA.
ID Q98R52
AC Q98R52
DI 01-NOV-1996 (FIREBASEL 01, Created)
DI 01-NOV-1996 (FIREBASEL 01, Last sequence update)
DI 01-NOV-1996 (FIREBASEL 01, Last annotation update)
DE glycoprotein precursor (Fragment).
OS Pseudorabies virus.
OC Viruses: DNA virus, no RNA stage: Herpesviridae:
OC Alphaherpesvirinae: Varicellovirinae.
OX NCBI TaxID 10545;
RN [11]

```

```

RP SEQUENCE FROM N.A.
RC STRAIN Becker.
RX MEDLINE=96070924; PubMed=7499261;
RA Ryan P., Edwards C.O.;
RT "Systematic introduction of proline in a eukaryotic signal sequence
RL J. Biol. Chem. 270:27876-27879(1995).
DR EMBL: U29125; AAC54545.1; -.
KW SIGNAL.
FT SIGNAL.
FT NON_TER.
SQ SEQUENCE 23 AA; 2271 MW; B7114E0B81406F3F CRC64;
  1 22 POTENTIAL.
  23 25
Query Match 33.3%; Score 22; DB 12; Length 23;
Best Local Similarity 62.5%; Pred. No. 8.5e+03;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 2 AKALKALL 9
DB 5 ARAMLALL 12

RESULT 32
Q69491
ID Q69491 PRELIMINARY; PRT; 23 AA.
AC Q69491;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE Glycoprotein C precursor (Fragment).
GN GC.
OS Pseudorabies virus.
OC Viruses; dsDNA viruses, no RNA stage; herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OX NCBI_TaxID:10345;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN Becker.
RX MEDLINE=95081163; PubMed=7989378;
RA Tomita M., Wilkinson F.S., Ryan P.,
RT "Can a signal sequence become too hydrophobic?";
RL J. Biol. Chem. 269:32016-32021(1994).
DR EMBL: L36969; AAA79966.1; -.
KW SIGNAL.
FT SIGNAL.
FT NON_TER.
SQ SEQUENCE 23 AA; 2299 MW; B711261E9146096F CRC64;
  1 22 POTENTIAL.
  23 24
Query Match 33.3%; Score 22; DB 12; Length 23;
Best Local Similarity 62.5%; Pred. No. 8.5e+03;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 2 AKALKALL 9
DB 5 ARAMLALL 12

RESULT 33
Q87084
ID Q87084 PRELIMINARY; PRT; 23 AA.
AC Q87084;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE Glycoprotein C precursor (Fragment).
GN GC.
OS Pseudorabies virus.
OC Viruses; dsDNA viruses, no RNA stage; herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OX NCBI_TaxID:10345;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN Becker.
RX MEDLINE=96070924; PubMed=7499261;
RA Ryan P., Edwards C.O.;
RT "Systematic introduction of proline in a eukaryotic signal sequence
RL J. Biol. Chem. 270:27876-27879(1995).
DR EMBL: U29125; AAC54545.1; -.
KW SIGNAL.
FT SIGNAL.
FT NON_TER.
SQ SEQUENCE 23 AA; 2271 MW; B7114E0B81406F3F CRC64;
  1 22 POTENTIAL.
  23 25
Query Match 33.3%; Score 22; DB 12; Length 23;
Best Local Similarity 62.5%; Pred. No. 8.5e+03;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 2 AKALKALL 9
DB 5 ARAMLALL 12

RESULT 34
Q8WYB9
ID Q8WYB9 PRELIMINARY; PRT; 24 AA.
AC Q8WYB9;
DT 01-MAR-2002 (TREMBLrel. 20, Created)
DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)
DT 01-MAR-2002 (TREMBLrel. 20, Last annotation update)
DE Dystrophin (Fragment).
GN DMO.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID:9606;
RN [1]
RP SEQUENCE OF 7-24 FROM N.A.
RX MEDLINE=97107974; PubMed 8950674;
RA Koest P.A., Bout M., van der Puijn A., Ginjaar L.B., Bakker E.,
RT Hoogerwerf F.B., van ommen G.J., den Dunnen J.L.;
RT "Splicing mutations in DMD/BBP detected by RT-PCR/RT: detection of a
RT 194A insertion in the cysteine rich domain of dystrophin (fragment)
RT with RMD.";
RL J. Med. Genet. 33:945-949(1996).
RN [2]
RP SEQUENCE FROM N.A.
RC Roberts R.G.;
RA Submitted (Nov-1999) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC den Dunnen J.L.;
RL Submitted (Dec-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF213441; AAL61585.1; -.
FT NON_TER.
FT NON_TER.
SQ SEQUENCE 24 AA; 2369 MW; AC93A264F4463E2 CRC64;
  1 24
Query Match 33.3%; Score 22; DB 4; Length 24;
Best Local Similarity 54.5%; Pred. No. 8.6e+03;
Matches 6; Conservative 1; Mismatches 4; Indels 0; Gaps 0;
QY 2 AKALKALL 12
DB 14 AMKLRRQKAL 24

RESULT 35
Q9T213
ID Q9T213 PRELIMINARY; PRT; 25 AA.
AC Q9T213;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE 16.1 kDa phosphosystem I PSAT Protein (Fragment).
OS Nicotiana sylvestris (Wood tobacco).

```

```

CG Chloroplast.
CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
CC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
CC Asterales; Lamiales; Solanales; Solanaceae; Nicotiana.
CX NCBI_TaxID 4096;
CN REV.
RP SEQUENCE.
RX MEDLINE 94105445; PubMed 8278548;
RA Okada J., Mikami K., Hayashida N., Nakamura M., Sugiura M.;
RT "Molecular heterogeneity of photosystem I, psal, psalE, psalF, psalH, and
RI psal are all present in isoforms in Nicotiana spp.;"
RL Plant Physiol. 102:1259-1267(1993);
DR InterPro: IP0003666; PSI_Psal;
DR Pfam: PF02507; PSI_Psal; 1;
SQ SEQUENCE 25 AA; 2844 MW; 26610FFELV22879E CRC64;

Query Match 33.4%; Score 22; DB P; Length 25;
Best Local Similarity 50.0%; Pred. No. 9.2e+04;
Matches 6; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 FAKALKALKAL 12
DB 1 1 1 1 1 1
14 FAKKKQXXXXK 25

RESULT 36
Q09072
AC Q09072 PRELIMINARY; PRT; 25 AA.
DI 01 NOV 1996 (TrEMBLrel. 01, Created)
DI 01 NOV 1996 (TrEMBLrel. 01, Last sequence update)
DI 01 MAR 2003 (TrEMBLrel. 23, Last annotation update)
DE P2 (Fragment).
CN P2.
CS Pinus radiata (Monterey pine).
CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
CC Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus.
CX NCBI_TaxID 3447;
CN REV.
RP SEQUENCE FROM N.A.
RA Kohn D.Y., Bradshaw P.E., Waller C., Cornett M., Pountain D.W.;
RT "Characterization of the MADS box gene family in Pinus radiata using
RI PCR cloning;"
RL Submitted (F01160) to EMBL, accession number F01160.
CC -1- SUBCELLULAR LOCATION: NUCLEUS (CYTOSOL).
CC -1- SIMILARITY: BELONGS TO THE MADS DOMAIN FAMILY OF TRANSSCRIPTION
CC FACTORS.
DR EMBL: U48626; AAA93470.1; 1;
DR InterPro: IPR002100; TF_MADSbox.
DR Pfam: PF00419; SRF_TF; 1;
DR PROSITE: PS50066; MADS_BOX_2; 1;
KW DNA-binding; Nuclear protein; Transcription; Transcription regulation.
FI NON_TER 1;
FI NON_TER 25; 25
SQ SEQUENCE 25 AA; 2880 MW; 020F92F5AF0CD5F9 CRC64;

Query Match 33.4%; Score 22; DB P; Length 25;
Best Local Similarity 50.0%; Pred. No. 9.2e+04;
Matches 6; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 FAKALKALKAL 10
DB 1 1 1 1 1 1
6 FSKKKKKL 14

RESULT 37
Q09072
AC Q09072 PRELIMINARY; PRT; 25 AA.
DI 01 JUN 2001 (TrEMBLrel. 17, Created)
DI 01 JUN 2001 (TrEMBLrel. 17, Last sequence update)
DI 01 OCT 2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment).
CN REV.

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```

CN REV.
CC Human immunodeficiency virus 1.
CC Viruses; Retroviruses; Retroviridae; Lentivirinae.
CX NCBI_TaxID 11790;
CN REV.
RP SEQUENCE FROM N.A.
RA STANIN IV000 17;
RX MEDLINE 21422926; PubMed 11429018;
RA Striba L.D., Trounstein F.K., Zeller M., Roush M.W.;
RT "Characterization and phylogenetic analysis of a novel accessory gene,
RI ALOS, from human retroviruses 17, 22, and 24 (2001)
DE EMBL: AF225522; AA09150.1;
DR InterPro: IPR000625; REV_Protein.
DR Pfam: PF00424; REV; 1;
FI NON_TER 25; 25
SQ SEQUENCE 25 AA; 2735 MW; 4F08F43134F134F6 CRC64;

Query Match 33.4%; Score 22; DB P; Length 25;
Best Local Similarity 40.0%; Pred. No. 9.2e+04;
Matches 4; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALIKALKAL 15
DB 1 1 1 1 1 1 1
10 FALQAVRLL 19

RESULT 38
Q09080
AC Q09080 PRELIMINARY; PRT; 25 AA.
DI 01 JUN 2001 (TrEMBLrel. 17, Created)
DI 01 JUN 2001 (TrEMBLrel. 17, Last sequence update)
DI 01 OCT 2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment).
CN REV.
CC Human immunodeficiency virus 1.
CC Viruses; Retroviruses; Retroviridae; Lentivirinae.
CX NCBI_TaxID 11790;
CN REV.
RP SEQUENCE FROM N.A.
RA STANIN IV013 3;
RX MEDLINE 21422927; PubMed 11429019;
RA Striba L.D., Trounstein F.K., Zeller M., Roush M.W.;
RT "Characterization and phylogenetic analysis of a novel accessory gene,
RI ALOS, from human retroviruses 17, 22, and 24 (2001)
DE EMBL: AF225526; AA09170.1;
DR InterPro: IPR000625; REV_Protein.
DR Pfam: PF00424; REV; 1;
FI NON_TER 25; 25
SQ SEQUENCE 25 AA; 2735 MW; 4F08F43134F134F6 CRC64;

Query Match 33.4%; Score 22; DB P; Length 25;
Best Local Similarity 40.0%; Pred. No. 9.2e+04;
Matches 4; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALIKALKAL 15
DB 1 1 1 1 1 1 1
10 FALQAVRLL 19

RESULT 39
Q09081
AC Q09081 PRELIMINARY; PRT; 25 AA.
DI 01 JUN 2001 (TrEMBLrel. 17, Created)
DI 01 JUN 2001 (TrEMBLrel. 17, Last sequence update)
DI 01 OCT 2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment).
CN REV.

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us-09-820-053a-43.rspt

Thu Aug 21 08:36:30 2003

```

OC Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09200.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

RESULT 40
Q99RQ0
ID Q99RQ0 PRELIMINARY; PRI: 25 AA.
AC Q99RQ0;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-oct-2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment)
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09145.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

RESULT 41
Q99R15
ID Q99R15 PRELIMINARY; PRI: 25 AA.
AC Q99R15;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-oct-2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment)
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09145.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

RESULT 42
Q99R19
ID Q99R19 PRELIMINARY; PRI: 25 AA.
AC Q99R19;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-oct-2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment)
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09145.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

RESULT 43
Q99R16
ID Q99R16 PRELIMINARY; PRI: 25 AA.
AC Q99R16;
DT 01-JUN-1999 (TrEMBLrel. 12, Created)
DT 01-JUN-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-oct-2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment)
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09145.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

RESULT 44
Q99R17
ID Q99R17 PRELIMINARY; PRI: 25 AA.
AC Q99R17;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-oct-2002 (TrEMBLrel. 22, Last annotation update)
DE Rev protein (Fragment)
GN REV.
OS Human immunodeficiency virus 1.
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.
OX NCBI_TaxID:11676;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN: IV001-2;
RX MEDLINE: 2142026; PubMed: 11429118;
RA Scriba T.J., Treurnicht F.K., Zeller M., Engelbrecht S.,
RA van Kousburg E.J.
RA "Characterization and phylogenetic analysis of South African HIV-1
RT subtype C accessory genes."
RL AIDS Res. Hum. Retroviruses 17:775-781(2001).
DR EMBL: AF425762; AAK09145.1;
DR InterPro: IPR000625; REV_protein.
DR Plam: PF00424; REV: 1.
DR NON_TER 25
SQ SEQUENCE 25 AA; 2735 MW; 4E934E445E4F224B CRC64;

Query Match 33.4%; Score 22; DB 15; Length 25;
Best Local Similarity 40.0%; Prod. No. 9.2e+03;
Matches 4; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 6 KALLKALKAL 15
DB 10 EALLQAVR11 19
|||||:
10 EALLQAVR11 19

```


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OM protein - protein search, using sw model

Run on: August 21, 2003, 08:14:25 : Search time 23 seconds
(without alignments)
30.670 Million cell updates/sec

Title: us-09-820-053a-43
Perfect score: 66
Sequence: 1 FAKALKALKALKAL 15

Scoring table: BLOSUM62

Gapop 10.0 : Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 1617

Minimum DB seq length: 0

Maximum DB seq length: 25

Post processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARYS

Result No.	Score	Query Match %	Length	EB	ID	Description
1	26	39.4	14	1	MCRX_METTM	P58815 methanobact
2	25	37.9	14	1	MAST_VESCR	P01516 vespa crabr
3	25	37.9	14	1	MAST_VESLE	P01514 vespa lew
4	25	37.9	14	1	MAST_VESMA	P04205 vespa ma
5	25	37.9	20	1	PCW6_PACGO	P82428 pachycondyl
6	25	37.9	21	1	BOH4_BOMVA	P91505 bombia var
7	24	36.4	14	1	MAST_VESOP	P17238 vespa orien
8	24	34.8	13	1	CHRD_PAPID	P42718 parapolylid
9	23	34.8	20	1	VIR1_VACGV	P15714 vaccinia vi
10	22	33.3	13	1	RS19_ASHYP	Q44592 ash yellow
11	22	33.3	21	1	JAP2_PANJA	P83706 raba japa
12	22	33.3	24	1	PCL1_PACGO	P82421 pachycondyl
13	22	33.3	21	1	PCL2_PACGO	P82422 pachycondyl
14	22	33.3	24	1	PG2_XHRLA	P39080 xenopus lac
15	21	31.8	16	1	ALRX_PSEPU	P17916 pseudomonas
16	21	31.8	25	1	V194_ARCFB	G30845 archaebactob
17	20	30.3	17	1	UP36_UPEMI	P82043 upercicola m
18	20	30.3	18	1	OBP_LYMDI	P34173 lymantria d
19	20	30.3	20	1	JHB_POMMO	P81627 bombyx mori
20	20	30.3	20	1	SYR_PAT	P40329 rattus norv
21	20	30.3	21	1	BOH1_BOMVA	P82282 bombyx mori
22	20	30.3	21	1	BOH4_BOMVA	P82284 bombyx mori
23	20	30.3	21	1	MDH_KLEPN	P80535 klebsiella
24	20	30.3	21	1	B523_SEREL	P83378 serratia pl
25	20	30.3	22	1	MLP_FANTC	P56924 tana tempor
26	20	30.3	22	1	PSL_PSEPD	P83188 pseudis par
27	19	28.8	24	1	TEMP_PAFTE	P56923 tana tempor
28	19	28.8	10	1	MAST_VESXA	P01515 vespa xanth
29	19	28.8	14	1	UP25_UPENI	P82031 upercicola i
30	19	28.8	19	1	MAX8_BOMMX	P83087 bombyx mori
31	19	28.8	20	1	RT16_BOVIN	P82015 bos taurus
32	19	28.8	20	1	RT16_BOVIN	P82015 bos taurus
33	19	28.8	24	1	FRMA_LITIN	P82023 titicola i

RESULT 1

ID	MCRX_METTM	STANDARD:	PRF:	14 AA.
AC	P58815;			
DT	28 FEB-2003 (Rel. 41, Last sequence update)			
DT	28 FEB-2003 (Rel. 41, Last sequence update)			
DE	Methyl-coenzyme M reductase 11 alpha subunit (DS 1.33.1.1) (90% id)			
DE	alpha) (Fragment).			
GR	META.			
OS	Methanobacterium thermoautotrophicum (strain Marburg / DSM 2133).			
OC	Archaea: Euryarchaeota: Methanobacteriales: Methanobacteriales:			
OC	Methanobacteriales: Methanobacteriales:			
OX	NCBI_TaxID:79929;			
RN	[1]			
RP	SEQUENCE.			
KX	RESIDUE 91079370; Pubmed 2269306;			
KA	Rospert S., Linder D., Eilermann J., Thauer R.K.:			
RT	"Two genetically distinct methyl-coenzyme M reductases in			
RT	Methanobacterium thermoautotrophicum strain Marburg and Delta H.:"			
KL	Eur. J. Biochem. 194:871-877(1990).			
CC	1. FUNCTION: Reduction of methyl-coenzyme M (2-(methylthio)ethanesulfonic acid) with 7-mercaptoheptanoylthioamine phosphate to methane and an heterodisulfide.			
CC	-1- CATALYTIC ACTIVITY: CH(3)-S-CoM + H-S-HP = CH(4) + CoM-S-S-HP.			
CC	1- COFACTOR: THE ENZYME COMPLEX BINDS TIGHTLY (NOT K+ COVALENTLY) TO TWO MOLECULES OF SERINE. THE 14-21-11-12-13-14-15-16-17-18-19-20-21-22-23-24-25-26-27-28-29-30-31-32-33-34-35-36-37-38-39-40-41-42-43-44-45-46-47-48-49-50-51-52-53-54-55-56-57-58-59-60-61-62-63-64-65-66-67-68-69-70-71-72-73-74-75-76-77-78-79-80-81-82-83-84-85-86-87-88-89-90-91-92-93-94-95-96-97-98-99-100-101-102-103-104-105-106-107-108-109-110-111-112-113-114-115-116-117-118-119-120-121-122-123-124-125-126-127-128-129-130-131-132-133-134-135-136-137-138-139-140-141-142-143-144-145-146-147-148-149-150-151-152-153-154-155-156-157-158-159-160-161-162-163-164-165-166-167-168-169-170-171-172-173-174-175-176-177-178-179-180-181-182-183-184-185-186-187-188-189-190-191-192-193-194-195-196-197-198-199-200-201-202-203-204-205-206-207-208-209-210-211-212-213-214-215-216-217-218-219-220-221-222-223-224-225-226-227-228-229-230-231-232-233-234-235-236-237-238-239-240-241-242-243-244-245-246-247-248-249-250-251-252-253-254-255-256-257-258-259-260-261-262-263-264-265-266-267-268-269-270-271-272-273-274-275-276-277-278-279-280-281-282-283-284-285-286-287-288-289-290-291-292-293-294-295-296-297-298-299-300-301-302-303-304-305-306-307-308-309-310-311-312-313-314-315-316-317-318-319-320-321-322-323-324-325-326-327-328-329-330-331-332-333-334-335-336-337-338-339-340-341-342-343-344-345-346-347-348-349-350-351-352-353-354-355-356-357-358-359-360-361-362-363-364-365-366-367-368-369-370-371-372-373-374-375-376-377-378-379-380-381-382-383-384-385-386-387-388-389-390-391-392-393-394-395-396-397-398-399-400-401-402-403-404-405-406-407-408-409-410-411-412-413-414-415-416-417-418-419-420-421-422-423-424-425-426-427-428-429-430-431-432-433-434-435-436-437-438-439-440-441-442-443-444-445-446-447-448-449-450-451-452-453-454-455-456-457-458-459-460-461-462-463-464-465-466-467-468-469-470-471-472-473-474-475-476-477-478-479-480-481-482-483-484-485-486-487-488-489-490-491-492-493-494-495-496-497-498-499-500-501-502-503-504-505-506-507-508-509-510-511-512-513-514-515-516-517-518-519-520-521-522-523-524-525-526-527-528-529-530-531-532-533-534-535-536-537-538-539-540-541-542-543-544-545-546-547-548-549-550-551-552-553-554-555-556-557-558-559-560-561-562-563-564-565-566-567-568-569-570-571-572-573-574-575-576-577-578-579-580-581-582-583-584-585-586-587-588-589-590-591-592-593-594-595-596-597-598-599-600-601-602-603-604-605-606-607-608-609-610-611-612-613-614-615-616-617-618-619-620-621-622-623-624-625-626-627-628-629-630-631-632-633-634-635-636-637-638-639-640-641-642-643-644-645-646-647-648-649-650-651-652-653-654-655-656-657-658-659-660-661-662-663-664-665-666-667-668-669-670-671-672-673-674-675-676-677-678-679-680-681-682-683-684-685-686-687-688-689-690-691-692-693-694-695-696-697-698-699-700-701-702-703-704-705-706-707-708-709-710-711-712-713-714-715-716-717-718-719-720-721-722-723-724-725-726-727-728-729-730-731-732-733-734-735-736-737-738-739-740-741-742-743-744-745-746-747-748-749-750-751-752-753-754-755-756-757-758-759-760-761-762-763-764-765-766-767-768-769-770-771-772-773-774-775-776-777-778-779-780-781-782-783-784-785-786-787-788-789-790-791-792-793-794-795-796-797-798-799-800-801-802-803-804-805-806-807-808-809-810-811-812-813-814-815-816-817-818-819-820-821-822-823-824-825-826-827-828-829-830-831-832-833-834-835-836-837-838-839-840-841-842-843-844-845-846-847-848-849-850-851-852-853-854-855-856-857-858-859-860-861-862-863-864-865-866-867-868-869-870-871-872-873-874-875-876-877-878-879-880-881-882-883-884-885-886-887-888-889-890-891-892-893-894-895-896-897-898-899-900-901-902-903-904-905-906-907-908-909-910-911-912-913-914-915-916-917-918-919-920-921-922-923-924-925-926-927-928-929-930-931-932-933-934-935-936-937-938-939-940-941-942-943-944-945-946-947-948-949-950-951-952-953-954-955-956-957-958-959-960-961-962-963-964-965-966-967-968-969-970-971-972-973-974-975-976-977-978-979-980-981-982-983-984-985-986-987-988-989-990-991-992-993-994-995-996-997-998-999-1000-1001-1002-1003-1004-1005-1006-1007-1008-1009-1010-1011-1012-1013-1014-1015-1016-1017-1018-1019-1020-1021-1022-1023-1024-1025-1026-1027-1028-1029-1030-1031-1032-1033-1034-1035-1036-1037-1038-1039-1040-1041-1042-1043-1044-1045-1046-1047-1048-1049-1050-1051-1052-1053-1054-1055-1056-1057-1058-1059-1060-1061-1062-1063-1064-1065-1066-1067-1068-1069-1070-1071-1072-1073-1074-1075-1076-1077-1078-1079-1080-1081-1082-1083-1084-1085-1086-1087-1088-1089-1090-1091-1092-1093-1094-1095-1096-1097-1098-1099-1100-1101-1102-1103-1104-1105-1106-1107-1108-1109-1110-1111-1112-1113-1114-1115-1116-1117-1118-1119-1120-1121-1122-1123-1124-1125-1126-1127-1128-1129-1130-1131-1132-1133-1134-1135-1136-1137-1138-1139-1140-1141-1142-1143-1144-1145-1146-1147-1148-1149-1150-1151-1152-1153-1154-1155-1156-1157-1158-1159-1160-1161-1162-1163-1164-1165-1166-1167-1168-1169-1170-1171-1172-1173-1174-1175-1176-1177-1178-1179-1180-1181-1182-1183-1184-1185-1186-1187-1188-1189-1190-1191-1192-1193-1194-1195-1196-1197-1198-1199-1200-1201-1202-1203-1204-1205-1206-1207-1208-1209-1210-1211-1212-1213-1214-1215-1216-1217-1218-1219-1220-1221-1222-1223-1224-1225-1226-1227-1228-1229-1230-1231-1232-1233-1234-1235-1236-1237-1238-1239-1240-1241-1242-1243-1244-1245-1246-1247-1248-1249-1250-1251-1252-1253-1254-1255-1256-1257-1258-1259-1260-1261-1262-1263-1264-1265-1266-1267-1268-1269-1270-1271-1272-1273-1274-1275-1276-1277-1278-1279-1280-1281-1282-1283-1284-1285-1286-1287-1288-1289-1290-1291-1292-1293-1294-1295-1296-1297-1298-1299-1300-1301-1302-1303-1304-1305-1306-1307-1308-1309-1310-1311-1312-1313-1314-1315-1316-1317-1318-1319-1320-1321-1322-1323-1324-1325-1326-1327-1328-1329-1330-1331-1332-1333-1334-1335-1336-1337-1338-1339-1340-1341-1342-1343-1344-1345-1346-1347-1348-1349-1350-1351-1352-1353-1354-1355-1356-1357-1358-1359-1360-1361-1362-1363-1364-1365-1366-1367-1368-1369-1370-1371-1372-1373-1374-1375-1376-1377-1378-1379-1380-1381-1382-1383-1384-1385-1386-1387-1388-1389-1390-1391-1392-1393-1394-1395-1396-1397-1398-1399-1400-1401-1402-1403-1404-1405-1406-1407-1408-1409-1410-1411-1412-1413-1414-1415-1416-1417-1418-1419-1420-1421-1422-1423-1424-1425-1426-1427-1428-1429-1430-1431-1432-1433-1434-1435-1436-1437-1438-1439-1440-1441-1442-1443-1444-1445-1446-1447-1448-1449-1450-1451-1452-1453-1454-1455-1456-1457-1458-1459-1460-1461-1462-1463-1464-1465-1466-1467-1468-1469-1470-1471-1472-1473-1474-1475-1476-1477-1478-1479-1480-1481-1482-1483-1484-1485-1486-1487-1488-1489-1490-1491-1492-1493-1494-1495-1496-1497-1498-1499-1500-1501-1502-1503-1504-1505-1506-1507-1508-1509-1510-1511-1512-1513-1514-1515-1516-1517-1518-1519-1520-1521-1522-1523-1524-1525-1526-1527-1528-1529-1530-1531-1532-1533-1534-1535-1536-1537-1538-1539-1540-1541-1542-1543-1544-1545-1546-1547-1548-1549-1550-1551-1552-1553-1554-1555-1556-1557-1558-1559-1560-1561-1562-1563-1564-1565-1566-1567-1568-1569-1570-1571-1572-1573-1574-1575-1576-1577-1578-1579-1580-1581-1582-1583-1584-1585-1586-1587-1588-1589-1590-1591-1592-1593-1594-1595-1596-1597-1598-1599-1600-1601-1602-1603-1604-1605-1606-1607-1608-1609-1610-1611-1612-1613-1614-1615-1616-1617-1618-1619-1620-1621-1622-1623-1624-1625-1626-1627-1628-1629-1630-1631-1632-1633-1634-1635-1636-1637-1638-1639-1640-1641-1642-1643-1644-1645-1646-1647-1648-1649-1650-1651-1652-1653-1654-1655-1656-1657-1658-1659-1660-1661-1662-1663-1664-1665-1666-1667-1668-1669-1670-1671-1672-1673-1674-1675-1676-1677-1678-1679-1680-1681-1682-1683-1684-1685-1686-1687-1688-1689-1690-1691-1692-1693-1694-1695-1696-1697-1698-1699-1700-1701-1702-1703-1704-1705-1706-1707-1708-1709-1710-1711-1712-1713-1714-1715-1716-1717-1718-1719-1720-1721-1722-1723-1724-1725-1726-1727-1728-1729-1730-1731-1732-1733-1734-1735-1736-1737-1738-1739-1740-1741-1742-1743-1744-1745-1746-1747-1748-1749-1750-1751-1752-1753-1754-1755-1756-1757-1758-1759-1760-1761-1762-1763-1764-1765-1766-1767-1768-1769-1770-1771-1772-1773-1774-1775-1776-1777-1778-1779-1780-1781-1782-1783-1784-1785-1786-1787-1788-1789-1790-1791-1792-1793-1794-1795-1796-1797-1798-1799-1800-1801-1802-1803-1804-1805-1806-1807-1808-1809-1810-1811-1812-1813-1814-1815-1816-1817-1818-1819-1820-1821-1822-1823-1824-1825-1826-1827-1828-1829-1830-1831-1832-1833-1834-1835-1836-1837-1838-1839-1840-1841-1842-1843-1844-1845-1846-1847-1848-1849-1850-1851-1852-1853-1854-1855-1856-1857-1858-1859-1860-1861-1862-1863-1864-1865-1866-1867-1868-1869-1870-1871-1872-1873-1874-1875-1876-1877-1878-1879-1880-1881-1882-1883-1884-1885-1886-1887-1888-1889-1890-1891-1892-1893-1894-1895-1896-1897-1898-1899-1900-1901-1902-1903-1904-1905-1906-1907-1908-1909-1910-1911-1912-1913-1914-1915-1916-1917-1918-1919-1920-1921-1922-1923-1924-1925-1926-1927-1928-1929-1930-1931-1932-1933-1934-1935-1936-1937-1938-1939-1940-1941-1942-1943-1944-1945-1946-1947-1948-1949-1950-1951-1952-1953-1954-1955-1956-1957-1958-1959-1960-1961-1962-1963-1964-1965-1966-1967-1968-1969-1970-1971-1972-1973-1974-1975-1976-1977-1978-1979-1980-1981-1982-1983-1984-1985-1986-1987-1988-1989-1990-1991-1992-1993-1994-1995-1996-1997-1998-1999-2000-2001-2002-2003-2004-2005-2006-2007-2008-2009-2010-2011-2012-2013-2014-2015-2016-2017-2018-2019-2020-2021-2022-2023-2024-2025-2026-2027-2028-2029-2030-2031-2032-2033-2034-2035-2036-2037-2038-2039-2040-2041-2042-2043-2044-2045-2046-2047-2048-2049-2050-2051-2052-2053-2054-2055-2056-2057-2058-2059-2060-2061-2062-2063-2064-2065-2066-2067-2068-2069-2070-2071-2072-2073-2074-2075-2076-2077-2078-2079-2080-2081-2082-2083-2084-2085-2086-2087-2088-2089-2090-2091-2092-2093-2094-2095-2096-2097-2098-2099-2100-2101-2102-2103-2104-2105-2106-2107-2108-2109-2110-2111-2112-2113-2114-2115-2116-2117-2118-2119-2120-2121-2122-2123-2124-2125-2126-2127-2128-2129-2130-2131-2132-2133-2134-2135-2136-2137-2138-2139-2140-2141-2142-2143-2144-2145-2146-2147-2148-2149-2150-2151-2152-2153-2154-2155-2156-2157-2158-2159-2160-2161-2162-2163-2164-2165-2166-2167-2168-2169-2170-2171-2172-2173-2174-2175-2176-2177-2178-2179-2180-2181-2182-2183-2184-2185-2186-2187-2188-2189-2190-2191-2192-2193-2194-2195-2196-2197-2198-2199-2200-2201-2202-2203-2204-2205-2206-2207-2208-2209-2210-2211-2212-2213-2214-2215-2216-2217-2218-2219-2220-2221-2222-2223-2224-22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CC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Vespoidae;
 CC Vespidae; Vespinae; Vespa.
 CX NCBI_TaxID: 7445;
 KN 11
 RP SEQUENCE.
 R0 TISSUE: Venom;
 KA Ariolas A., Pisano J.J.;
 KA "Isolation and characterization of two new peptides, mastoparan C and
 RT crabrolin, from the venom of the European hornet, *Vespa crabro* L."
 RL J. Biol. Chem. 259:10106-10111(1984).
 CC 1- FUNCTION: Mast cell degranulating peptide. Activates G proteins
 CC that couple to phospholipase C.
 DR PIR: A01779; QMVP2.
 KW Mast cell degranulation; Amidation.
 FT MOD_RES 14 AA: AMIDATION.
 SQ SEQUENCE 14 AA: 1508 MW: 5500CECA1D6AR1D7 CRC64:

Query Match 37.9% Score 25; DB 1; Length 14;
 Best Local Similarity 54.5% Pred. No. 3,3e+02;
 Matches 6; Conserved 1; Mismatches 4; Indels 0; Gaps 0;

QY 5 LKALLKALKAL 15
 11111111
 DB 3 LKALLAVAKK 13

RESULT 4

MAST_VESLE
 ID MAST_VESLE STANDARD; PRT: 14 AA.
 AC P01514;
 DT 21 JUL 1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Mastoparan.
 CC Vespa Lewisii (Yellow Jacket) (Wasp).
 CC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Vespoidae;
 CC Vespidae; Vespinae; Vespa.
 CX NCBI_TaxID: 7452;
 KN 11
 RP SEQUENCE, AND SYNTHESIS.
 R0 TISSUE: Venom;
 KA Hirai Y., Yasuhara T., Yoshida H., Nakajima H., Kitada G.;
 KA "A new mast cell degranulating peptide 'mastoparan' in the venom of
 RT Vespa Lewisii";
 RL Chem. Pharm. Bull. 27:1942-1944(1979).
 CC 1- FUNCTION: Mast cell degranulating peptide. Activates G proteins
 CC that couple to phospholipase C.
 DR PIR: A01776; QMVAV.
 DR PIR: 1979L; 20-SEP-81.
 KW Mast cell degranulation; Amidation; 3D-structure.
 FT MOD_RES 14 AA: 14 AMIDATION.
 SQ SEQUENCE 14 AA: 1480 MW: 6500CECA1D7R000 CRC64:

Query Match 37.9% Score 25; DB 1; Length 14;
 Best Local Similarity 60.0% Pred. No. 4,3e+02;
 Matches 6; Conserved 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 KATKALKAL 12
 11111111
 DB 4 KATKAVAKK 13

RESULT 4

MAST_VESMA
 ID MAST_VESMA STANDARD; PRT: 14 AA.
 AC P04205;
 DT 20-MAR-1987 (Rel. 04, Created)
 DT 20-MAR-1987 (Rel. 04, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Mastoparan M (Mast cell-degranulating peptide).

CC Vespa mandarinia (Hornet).
 CC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Vespoidae;
 CC Vespidae; Vespinae; Vespa.
 CX NCBI_TaxID: 7446;
 KN 11
 RP SEQUENCE.
 R0 TISSUE: Venom;
 KA Hirai Y., Yasuhara T., Yoshida H., Nakajima H.;
 KA "A new mast cell degranulating peptide, mastoparan M, in the venom
 RT the hornet *Vespa mandarinia*."
 RL Biochem. Res. 2:447-449(1981).
 CC 1- FUNCTION: Mast cell degranulating peptide. Activates G proteins
 CC that couple to phospholipase C.
 DR PIR: A01777; QMVMH.
 KW Mast cell degranulation; Amidation.
 FT MOD_RES 14 AA: AMIDATION.
 SQ SEQUENCE 14 AA: 1480 MW: 6560A09A1D7R000 CRC64:

Query Match 37.9% Score 25; DB 1; Length 14;
 Best Local Similarity 60.0% Pred. No. 4,3e+02;
 Matches 6; Conserved 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 KATKALKAL 12
 11111111
 DB 4 KATKAVAKK 13

RESULT 5

MAST_PACGO
 ID MAST_PACGO STANDARD; PRT: 20 AA.
 AC P82428;
 DT 16 OCT 2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Ponerine W6.
 CC Pachycondyla gonidii (Ponerine ant).
 CC Pachycondyla; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Eumecoptera;
 CC Ponerinae; Pachycondylina.
 CX NCBI_TaxID: 110606;
 KN 11
 RP SEQUENCE, FUNCTION, AND AMIDATION.
 R0 TISSUE: Venom;
 EX MEDLINE 21294442; P01254 (1/9/90);
 KA Grivel J., Becker V., Le Goff G., Poirier F., Krieger T., Bouček A., Poirier
 KA Lomonte A., Chabotier A., Lejeune A., Posselt J.;
 KA "Ponerine, new antibacterial and insecticidal peptides from the
 RT venom of the ant *Pachycondyla gonidii* W."
 RL J. Biol. Chem. 276:17823-17829(2001).
 CC 1- FUNCTION: ACTIVITY AGAINST SEVER POSITIVE INACTIVE INHIBITORS.
 CC 1- SUBCELLULAR LOCATION: Secreted.
 CC 1- MASS SPECIFICITY: MW 2029.40; METHIONINE-MODIFIED.
 KW Antibiotic; Insect immunity; Remolysing; Amidation.
 FT MOD_RES 20 AA: 20 AMIDATION.
 SQ SEQUENCE 20 AA: 2030 MW: 5241E1620A0A000 CRC64:

Query Match 37.9% Score 25; DB 1; Length 20;
 Best Local Similarity 41.7% Pred. No. 4,4e+02;
 Matches 6; Conserved 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 AKATKALKAL 13
 11111111
 DB 9 ASATKAVAKK 20

RESULT 6

MAST_PACMA
 ID MAST_PACMA STANDARD; PRT: 23 AA.
 AC P01505;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)

```

DT 15 SEP-2003 (Rel. 42, Last annotation update)
DE Bombinin.
OS Bombina variegata (Yellow-bellied toad).
OC Fukuyofa; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Archeobatrachia; Bombinatoridae; Bombina.
OX NCBI_TaxID-8448;
RN 111
RP SEQUENCE.
RC TISSUE: Skin secretion;
RA Csordas A., Michl H.;
RT "Isolation and structural resolution of a haemolytically active
RL polypeptide from the immune secretion of a European toad.";
RL Monatsh. Chem. 101:182-189(1970).
CC -1- FUNCTION: Has antimicrobial and hemolytic activities.
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- TISSUE SPECIFICITY: Skin.
CC -1- SIMILARITY: BELONGS TO THE BOMBININ FAMILY.
DR PIR: A01766; HMTD.
FW Amphibian defense peptide. Antibioidic; hemolysis; Amidation.
FT MOD_RES 24 AA: 2294 MW: ACCDFCEB7402E95 CRC64;
SQ SEQUENCE 24 AA: 2294 MW: ACCDFCEB7402E95 CRC64;

Query Match 37.9%; Score 25; DB 1; Length 24;
Best Local Similarity 62.7%; Pred. No. 5,5e+02;
Matches 6; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 ALKALKAL 12
DB 10 ALKGLAKGL 18

RESULT 7
MAST_VESOR STANDARD; PRT; 14 AA.
AC P17238;
DT 01-AUG-1990 (Rel. 15, Created)
DI 01-AUG-1990 (Rel. 15, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DE Mustoparan (Histamine releasing peptide 1) (HR-1).
OS Vespa orientalis (Oriental hornet).
OC Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Vespoidea;
OC Vespidae; Vespinae; Vespa.
OX NCBI_TaxID-7447;
RN 111
RP SEQUENCE.
RC TISSUE: Venom;
RA Miroshnikov A. I., Stachurski L.S.;
RA Rozynov B.V., Gushchin I.S.;
RT "Structure and properties of histamine releasing peptides from the
RI venom of Vespa orientalis hornet.";
RI Bioorg. Khim. 7:1467-1477(1981).
CC -1- FUNCTION: Mast cell desgranulating peptide. Activates G proteins
CC that couple to phospholipase C.
DR PIR: JN0489; JN0489.
KW Mast cell desgranulation; Amidation.
FT MOD_RES 14 AA: 1494 MW: CB4F9ECA026B00DD CRC64;
SQ SEQUENCE 14 AA: 1494 MW: CB4F9ECA026B00DD CRC64;

Query Match 36.4%; Score 24; DB 1; Length 14;
Best Local Similarity 62.5%; Pred. No. 4,8e+02;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 4 KALKALK 10
DB 4 KATAALK 11

RESULT 8
CHEP_PAKID STANDARD; PRT; 13 AA.
AC I42718;
DT 01-NOV-1995 (Rel. 32, Created)

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DT 01-NOV-1995 (Rel. 32, Last sequence update)
DI 01-NOV-1995 (Rel. 32, Last annotation update)
DE Chemotactic peptide.
OS Parapolybia indica.
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata; Vespoidea;
OC Vespidae; Polistinae; Parapolybia.
OX NCBI_TaxID-31921;
RN 111
RP SEQUENCE.
RC TISSUE: Venom;
RA Toki T., Yasuhara T., Nakajima T.;
RT "Isolation and sequential analysis of peptides on the venom sac of
RL Parapolybia indica.";
RL Eisai Dobutsu 39:105-111(1988).
KW Chemotaxis; Amidation.
FT MOD_RES 13 AA: 1298 MW: 56950; ERF39D9873 CRC94;
SQ SEQUENCE 13 AA: 1298 MW: 56950; ERF39D9873 CRC94;

Query Match 34.8%; Score 23; DB 1; Length 13;
Best Local Similarity 62.5%; Pred. No. 5,5e+02;
Matches 5; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 8 LKALKAL 15
DB 4 LKGLSAL 11

RESULT 9
V101_VACV STANDARD; PRT; 20 AA.
AC P16714;
DT 01-AUG-1990 (Rel. 15, Created)
DI 01-AUG-1990 (Rel. 15, Last sequence update)
DE 16-OCT-2001 (Rel. 40, Last annotation update)
DE Protein 11 (Fragment).
GN 111.
OS Vaccinia virus (strain WR).
OC Viruses; dsDNA viruses, no RNA stage; Poxviridae; Chordopoxvirinae;
OC Orthopoxvirus.
OC NCBI_TaxID-10254;
RN 111
RP SEQUENCE FROM N.A.
RX MEDLINE-88215015; PubMed-2835495;
RA Schmitt J.F.C., Stamenberg H.G.;
RT "Sequence and transcriptional analysis of the Vaccinia virus HindIII
RI 1 fragment.";
RL J. Virol. 62:1889-1897(1988).
CC -1- SIMILARITY: BELONGS TO THE POXVIRUS-11 FAMILY.
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EMBL: J04399; AAR59803.1;
DR InterPro; IPR004965; Pfam_11.
DR Pfam; PF04289; Pfam_11;
KW Late protein.
FT NON_TER 20 20
SQ SEQUENCE 20 AA: 2241 MW: 6197AA7866664B1 CRC94;

Query Match 34.8%; Score 23; DB 1; Length 20;
Best Local Similarity 83.4%; Pred. No. 2,6e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 AVALKA 7
DB 15 AVALKA 20

```


RT "Jenotricins, new antibacterial and insecticidal peptides from the
 RL venom of the ant *Pachycondyla goeldii*.";
 CC J. Biol. Chem. 276:17822-17829(2001).
 CC -f- FUNCTION: BROAD SPECTRUM OF ACTIVITY AGAINST BOTH GRAM-POSITIVE
 CC AND GRAM-NEGATIVE BACTERIA. HAS NON-HEMOLYTIC ACTIVITY.
 CC -f- SUBCELLULAR LOCATION: Secreted.
 CC -f- MASS SPECTROMETRY: MW:2576.67; METHOD:MALDI.
 KW Antibiotic; Insect immunity; Amidation.
 FT MOD_RES 24 24
 SQ SEQUENCE 24 AA; 2578 MW; 37830D5761515E8F CRC64;

Query Match 34.3%; Score 22; DB 1; Length 24;
 Best Local Similarity 50.0%; Pred. No. 1.6e+03;
 Matches 5; Conservations 2; Mismatches 3; Indels 0; Gaps 0;

QY 6 KALLIKALIN 15
 DB 14 KALLIKALIN 24

RESULT 14
 ID PQQ_XENIA STANDARD; PRT: 24 AA.
 AC P30080;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 15-FEB-1995 (Rel. 31, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Antimicrobial peptide PQQ.
 OS Xenopus laevis (African clawed frog).
 CC Enkaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidea; Pipidae;
 CC Xenopodidae; Xenopus.
 CC NCBI_TaxID=8355;
 RN [1]
 RP TISSUE:Stomach;
 RC SEQUENCE.

Query Match 33.4%; Score 22; DB 1; Length 24;
 Best Local Similarity 71.4%; Pred. No. 1.6e+03;
 Matches 5; Conservations 1; Mismatches 1; Indels 0; Gaps 0;

QY 4 ALKALIK 10
 DB 17 ALKALIK 24

RESULT 15
 ID ALRX_PSEPI STANDARD; PRT: 16 AA.
 AC P17916;
 DT 01-AUG-1990 (Rel. 15, Created)
 DT 01-AUG-1990 (Rel. 15, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Broad-spectrum antimicrobial peptide.
 OS Pseudomonas putida.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 CC Pseudomonadaceae; Pseudomonas.

OX NCBI_TaxID=303;
 RN [1]
 RP SEQUENCE.
 RC STRAIN AKU 0813;
 CC MEDLINE 85072810; PubMed-64 19247;
 RA "Inactivation of the Pseudomonas striata broad specificity aminopeptidase
 RT by D and L isomers of beta substituted amines: Kinetics,
 RL stoichiometry, active site peptide, and mechanistic studies.";
 CC -f- BIOCHEMISTRY 24:5195-5201(1984).
 CC -f- COFACTOR: Pyridoxal phosphate.
 CC -f- SIMILARITY: Belongs to the alanine racemase family.
 DR A29723; A29720.
 DR HAMAP: MF 01201; atypical; 1.
 DR InterPro: IPR000821; Ala_racemase.
 DR PROSITE: PS00395; ALANINE_RACEMASE; 1.
 KW isomerase; Pyridoxal phosphate.
 FT NON_TER 1 1
 FT AC1_SITE 6 6
 FT BINDING 6 6
 FT NCBI_TER 16 16
 SQ SEQUENCE 16 AA; 1572 MW; 9C6C25A11342E907 CRC64;

Query Match 31.8%; Score 21; DB 1; Length 16;
 Best Local Similarity 71.4%; Pred. No. 1.6e+03;
 Matches 5; Conservations 1; Mismatches 1; Indels 0; Gaps 0;

QY 5 LKALLKA 11
 DB 1 LKALLKA 7

RESULT 16
 ID Y194_ARCF9 STANDARD; PRT: 25 AA.
 AC Q30045;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hypothetical protein AF0194.
 CC Archaeobacteria; Archaeobacteria; Archaeobacteria;
 CC Archaeobacteria; Archaeobacteria;
 CC NCBI_TaxID=2234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN V.11 / DSM 4304 / AF0194;
 RX MEDLINE 9489475; PubMed 9489475;
 RA Kleck H., Clayton R.A., Tomb J.F., White O., Nelson K.L.,
 RA Richardson K.A., Dodson R.A., Tomb J.F., White O., Nelson K.L.,
 RA Richardson K.A., Dodson R.A., Tomb J.F., White O., Nelson K.L.,
 RA Fleischmann R.D., Quackenbush J., Lee N.A., Sutton G., Gillis S.,
 RA Kirkness E.F., Baugherty B.A., McInerney K., Adams M.L., Loftus B.,
 RA Peterson S., Reich C.L., McNeil L.K., Badger J.H., Shanks A.,
 RA Overbeek R., Gocayne J.D., Weidman J.F., McDaniel L., Olsen G.J.,
 RA Cotton M.D., Spillars T., Artlich P., Kaine R.P., Sykes S.M.,
 RA Sadow P.W., Anderson K.P., Bosman C., Fulton R., Starnali S.A.,
 RA Mason T.M., Olsen G.J., Fraser J.M., Smith H.O., Woese C.R.,
 RA Venter J.C.;
 RC "The complete genome sequence of the hyperthermophilic, sulphate-
 RT reducing archaeon *Archaeoglobus fulgidus*.";
 RL Nature 392:664-670(1997).

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DB EMBL: AF001094; AAB01054.1;
DB PIR: B69274; B69274.
DB TrEMBL: AF001094.1;
KW Hydrophobic protein; Complete proteome.
SQ SEQUENCE 25 AA: 2828 MW: 54380.60 467419F CRG64;

Query Match 31.8%; Score 21; DB 1; Length 25;
Best Local Similarity 57.1%; Prod. No. 2,466,043;
Matches 4; Conservative 2; Mismatches 1; Gaps 0;

QY 6 KALKAL 12
   1 1 1 1 1
DB 4 KAVIKGM 10

RESULT 17
DB UP36_UPEMJ STANDARD; PRI: 17 AA.
A* P82043;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Uperin 3.6.
OS Uperoleia mjoberti (Australian toadlet).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Myobatrachidae;
OC Myobatrachinae; Uperoleia.
CX NCBI TaxID 104964;
KN [1]
KW SEQUENCE, AND MASS SPECTROMETRY.
KO TISSUE SKIN SECRETION.
RA Bradford A.M., Bowle J.H., Taylor M.J., Wallace J.C.;
RT "New antitoxic uperin peptides from the dorsal glands of the
RL Aust. J. Chem. 49:1325-1341(1996).
CV 1 FUNCTION: SHOWS ANTIBACTERIAL ACTIVITY AGAINST BACTERIA, LACTIC
CV 1 LACTICACID, MOLDERS, SAGREES, S. FIDELMIS AND S. BERSIS.
CV 1 SUBCELLULAR LOCATION: Secreted.
CV 1 TISSUE SPECIFICITY: Expressed by the skin dorsal glands.
CV 1 MASS SPECTROMETRY: MW 1826; METHOD FAB.
KW Amphibian defense peptide; Antibiotic; Amidation.
PI M36-RES 17
SQ SEQUENCE 17 AA: 1778 MW: 7840884626 6A30 CRG64;
   1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
QY 4 ALKALKAL 15
DB 5 AAKVVNVUKNL 16

RESULT 18
DB UP36_UPEMJ STANDARD; PRI: 18 AA.
A* P34174;
DT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE General odorant-binding protein (G99) (Fragment).
OS Lymantiria dispar (Gypsy moth).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia; Noctuoidea;
OC Lymantiridae; Lymantiria.
CX NCBI TaxID 13124;
KN [1]
KW SEQUENCE.
KO MEDLINE 91186129; PubMed 2010751;
RA Vaut R.G., Prestwich G.D., Lerner M.R.;
RT "Odorant-binding protein subfamilies associate with distinct classes
RL J. Neurobiol. 22:74-84(1991).

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CV 1 FUNCTION: PRESENT IN THE ALD AND FLD FOLDING DOMAINS. MAY BE A
CV 1 SENSORY PROPERTIES AND ARE THROUGH 1. AL AND FLD DOMAINS. AN
CV 1 TRANSPEPTIDASE HYDROPHOBIC DOMAINS INTERACT WITH THE FLD
CV 1 SUBUNIT. Homodimer (Predicted)
CV 1 TISSUE SPECIFICITY: ANTENNA.
CV 1 SIMILARITY: RELATIONS TO THE PRO/5000 FAMILY
KW Collection Transport 18
PI NON-TER 18
SQ SEQUENCE 18 AA: 1994 MW: 8684808854 6A30
   1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
QY 1 FAKALK 6
DB 12 FAKALE 17

RESULT 19
DB UP36_UPEMJ STANDARD; PRI: 20 AA.
A* P81627;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Juvenile hormone binding protein (Fragment).
OS Bombyx mori (Silk moth).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia; Bombyx.
CX NCBI TaxID 7094;
KN [1]
KW SEQUENCE.
KO STRAIN BacKokJung TISSUE Bombyx;
RA Park C. H., Kim H.R.;
RT "Characterization of high affinity juvenile hormone binding protein
RL the Bombyx mori."
RL Journal of Insect Physiology 47:495-503(1994).
KN [2]
PI IDENTIFICATION OF CYS 9.
RA Park C. H.;
RL Submitted (09-1999) To The Swiss Prot Data Bank
CV 1 FUNCTION: PRESENT IN THE BOMBYX MORI FLD AND FLD DOMAINS.
CV 1 GENERAL INTERACTION BY BOMBYX MORI FLD AND FLD DOMAINS.
CV 1 SUBCELLULAR LOCATION: Secreted.
PI NON-TER 20
SQ SEQUENCE 20 AA: 2090 MW: 8669284842 6A30
   1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
QY 7 ALKALK 14
DB 4 ALKALK 16

RESULT 20
DB UP36_UPEMJ STANDARD; PRI: 21 AA.
A* P40479;
DT 01-FEB-1998 (Rel. 31, Created)
DT 01-FEB-1998 (Rel. 31, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Acetyl-L-tyrosine synthase (P40479) (Fragment).
OS RARS.
KN RARS.
KO RARS;
RA Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciuroidea; Muridae; Murinae; Murinae;
CX NCBI TaxID 10114;

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23 8 KALKAL 12
 14 11 11
 15 15-JUL-1998 (Rel. 46, created)
 16 15-JUL-1998 (Rel. 46, last sequence update)
 17 15-SEP-2003 (Rel. 42, last annotation update)
 18 SPECIES: *Litoria caerulea* (Green tree frog)
 19 LITOKIA SPLENDIDA (Mantidivert frog frog)
 20 EKATYOTA MOKATOA (Mantidivert frog frog)
 21 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 22 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 23 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 24 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 25 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 26 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 27 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 28 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 29 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 30 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 31 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 32 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 33 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 34 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 35 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 36 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 37 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 38 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 39 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 40 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 41 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 42 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 43 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 44 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 45 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 46 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 47 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 48 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 49 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 50 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)

51 8 KALKAL 15
 52 11 11
 53 15-JUL-1998 (Rel. 46, created)
 54 15-JUL-1998 (Rel. 46, last sequence update)
 55 15-SEP-2003 (Rel. 42, last annotation update)
 56 SPECIES: *Litoria caerulea* (Green tree frog)
 57 LITOKIA SPLENDIDA (Mantidivert frog frog)
 58 EKATYOTA MOKATOA (Mantidivert frog frog)
 59 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 60 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 61 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 62 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 63 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 64 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 65 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 66 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 67 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 68 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 69 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 70 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 71 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
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 78 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 79 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)
 80 AMPHIBIA MANTIDIVERT (Mantidivert frog frog)

RESULT 42

220554

hemocyanin subunit V - Atlantic horseshoe crab (Limulus)

C1Species: Limulus polyphemus (Atlantic horseshoe crab)

C1Date: 05-Jun-1997 #sequence_revision 05-Jun-1987 #text_change 08-Jul-1994

C1Accession: J20554

K1Name: J. J. Lamy, J. J. Sizer, P. V. Billiard, P. J. Jolles, P. J. Jolles, J. J. Feldmann, R. J.

Biochemistry 22: 5573-5585, 1988

A1Title: Quaternary structure of Limulus polyphemus hemocyanin.

A1Accession number: A00478

A1Molecule type: Protein

A1Residues: 1-24-1AM

C1Keywords: Limulus polyphemus hemocyanin is an association of eight different subunits

Query Match: 30.38; Score 20; DB 2; Length 24;

Best Local Similarity: 45.58; Pred. No. 5.7e+03;

Matches: 5; Conservative: 1; Mismatches: 5; Indels: 0; Gaps: 0;

QY 5 EKAL:KAKAL 15
1 1 1 1 1 1
16 KUKV:KUKV 12

RESULT 43

B04710

hemolymph protein HP1528 - Helicobacter pylori (strain 26695)

C1Species: Helicobacter pylori

C1Date: 09-Aug-1997 #sequence_revision 09-Aug-1997 #text_change 08-Oct-1999

C1Accession: H04710

K1Name: J. F. White, G. J. Kortvaag, A. R. Clayton, R. A. Sutton, G. G. Fleisemann, R. D.

Biochemistry 22: 5573-5585, 1988

A1Title: Study on N-terminal sequence of the haemoglobin of Volpes vulpes fox.

A1Accession number: P00617

A1Molecule type: Protein

A1Residues: 1-25-1M

C1Keywords: hemoglobin; globin; globin hemology

Query Match: 30.38; Score 20; DB 2; Length 25;

Best Local Similarity: 40.08; Pred. No. 5.9e+03;

Matches: 4; Conservative: 2; Mismatches: 4; Indels: 0; Gaps: 0;

QY 6 EKAL:KAKAL 15
1 1 1 1 1 1
16 KUKV:KUKV 25

RESULT 44

P00114

hemolymph beta chain - red fox (Lutra)

C1Species: Vulpes vulpes (red fox)

C1Date: 15-Jan-1994 #sequence_revision 15-Jan-1994 #text_change 19-May-2000

C1Accession: P00114

K1Name: J. J. Lamy, J. J. Sizer, P. V. Billiard, P. J. Jolles, P. J. Jolles, J. J. Feldmann, R. J.

Biochemistry 22: 5573-5585, 1988

A1Title: Study on N-terminal sequence of the haemoglobin of Volpes vulpes fox.

A1Accession number: P00114

A1Molecule type: Protein

A1Residues: 1-25-1M

C1Keywords: hemoglobin; globin; globin hemology

Query Match: 29.58; Score 19.5; DB 2; Length 15;

Best Local Similarity: 66.78; Pred. No. 4.3e+03;

Matches: 8; Conservative: 1; Mismatches: 2; Indels: 1; Gaps: 1;

QY 5 EKAL:KAKAL 15

db 3 EKAL:KAKAL 14

RESULT 45

A38841

rhodopsin homologue - squid (Loligo setacea) (Limulus)

C1Species: Loligo setacea (squid)

C1Date: 17-Jul-1992 #sequence_revision 17-Jul-1992 #text_change 01-Oct-1997

C1Accession: A38841

K1Name: M. J. Kubota, T. Hiraki, K. J. Kito, Y.

Biochemistry 22: 5573-5585, 1988

A1Title: Study on N-terminal sequence of the retinal binding site of squid visual pigment.

A1Accession number: P00617

A1Molecule type: Protein

A1Residues: 1-11-1M

C1Keywords: rhodopsin

C1Keywords: chromophore; retinal

C1Keywords: chromophore; retinal

Query Match: 28.88; Score 19; DB 2; Length 11;

Best Local Similarity: 100.08; Pred. No. 4.8e+03;

Matches: 4; Conservative: 0; Mismatches: 0; Indels: 0; Gaps: 0;

QY 1 EKAL:KAKAL 4
1 1 1 1
16 KUKV:KUKV 4

Search completed: August 21, 2003, 08:22:01

Job time: 42 secs

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Genetic version 5.1.6
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EM FASTA protein search, using SW model

Run on: August 21, 2003, 08:19:10 : Search time 55 Seconds

(without alignments)
 45,990 Million cell updates/sec

Filter: US-09-820-053A-43

Percent score: 62

Sequence: 1 PARAFACALFAALFAAL 15

Scoring table: RD-SIM62

Gap: 10.0, Gapext 0.5

Scoring: 44779 seqs, 131961718 residues

1st 10 number of hits satisfying chosen parameters: 111549

Minimum 1st seq length: 0

Maximum 1st seq length: 25

Post-processor: Minimum Match 0%

Listing first 45 summaries

Database:	Published_Applications-AA*
1	US07_Protein_Pep*
2	US07_Protein_Pep*
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45	US07_Protein_Pep*

Prod. No. 18: the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SWAPPAGE

1	US-09-820-053A-43	Sequence 43, Application No. US20010109171	15	100.0%	100.0%	15	0.0%
2	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
3	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
4	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
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19	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
20	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
21	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
22	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
23	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
24	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
25	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
26	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
27	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
28	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
29	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
30	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
31	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
32	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
33	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
34	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
35	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
36	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
37	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
38	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
39	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
40	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
41	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
42	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
43	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
44	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
45	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%

RESULTS

1	US-09-820-053A-43	Sequence 43, Application No. US20010109171	15	100.0%	100.0%	15	0.0%
2	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
3	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
4	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
5	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
6	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
7	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
8	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
9	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
10	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
11	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
12	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
13	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
14	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
15	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
16	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
17	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
18	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
19	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
20	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
21	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
22	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
23	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
24	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
25	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
26	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
27	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
28	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
29	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
30	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
31	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
32	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
33	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
34	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
35	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
36	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
37	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
38	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
39	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
40	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
41	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
42	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
43	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
44	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%
45	US-09-820-053A-43	Sequence 43, Application No. US20010109452A1	15	100.0%	100.0%	15	0.0%


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1  APPLICANT: OWEN, DONALD R.
2  TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE
3  FILE REFERENCE: HELIX028
4  CURRENT APPLICATION NUMBER: US/09/109,171
5  CURRENT FILING DATE: 2002-03-28
6  NUMBER OF SEQ ID NOS: 165
7  SOFTWARE: PATENTIN Ver. 2.1
8  SEQ ID NO: 13
9  LENGTH: 19
10 TYPE: PRT
11 ORGANISM: ARTIFICIAL SEQUENCE
12 FEATURE:
13 OTHER INFORMATION: SYNTHETIC SEQUENCE
14 US-10-109-171-13

Query Match      64.68; Score 42; DB 15; Length 19;
Best Local Similarity 78.98; Pred. No. 2.5;
Matches 11; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

CY 1 FAKAKALKALK 15
1 | | | | | | | | |
DB 1 FAKAKALKALK 15

RESULT 17
US-09-820-053A-68
Sequence 68; Application US/09/820053A
Publication No. US20030084243A1
GENERAL INFORMATION:
1 APPLICANT: OWEN, DONALD R.
2 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES
3 FILE REFERENCE: HELIX027
4 CURRENT APPLICATION NUMBER: US/09/820,053A
5 CURRENT FILING DATE: 2001-03-28
6 NUMBER OF SEQ ID NOS: 165
7 SOFTWARE: PATENTIN Ver. 2.1
8 SEQ ID NO: 68
9 LENGTH: 20
10 TYPE: PRT
11 ORGANISM: ARTIFICIAL SEQUENCE
12 FEATURE:
13 OTHER INFORMATION: SYNTHETIC SEQUENCE
14 NAME/KEY: MOD_PES
15 LOCATION: (203)
16 OTHER INFORMATION: AMIDATION
17 US-09-820-053A-68

Query Match      63.68; Score 42; DB 11; Length 20;
Best Local Similarity 76.98; Pred. No. 2.5;
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

CY 1 FAKAKALKALK 13
1 | | | | | | | | |
DB 5 FFAAKKFAKALF 17

RESULT 18
US-10-109-171-68
Sequence 68; Application US/10/109,171
Publication No. US20030109452A1
GENERAL INFORMATION:
1 APPLICANT: OWEN, DONALD R.
2 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE
3 FILE REFERENCE: HELIX028
4 CURRENT APPLICATION NUMBER: US/10/109,171
5 CURRENT FILING DATE: 2002-03-28
6 NUMBER OF SEQ ID NOS: 165
7 SOFTWARE: PATENTIN Ver. 2.1
8 SEQ ID NO: 68
9 LENGTH: 20
10 TYPE: PRT
11 ORGANISM: ARTIFICIAL SEQUENCE
12 FEATURE:
13 OTHER INFORMATION: SYNTHETIC SEQUENCE
14 NAME/KEY: MOD_PES
15 LOCATION: (203)
16 OTHER INFORMATION: AMIDATION
17 US-09-820-053A-68

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1  OTHER INFORMATION: SYNTHETIC SEQUENCE
2  NAME/KEY: MOD_PES
3  LOCATION: (20)
4  OTHER INFORMATION: AMIDATION
5  US-10-109-171-68

Query Match      64.68; Score 42; DB 15; Length 20;
Best Local Similarity 76.98; Pred. No. 2.5;
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

CY 1 FAKAKALKALK 13
1 | | | | | | | | |
DB 5 FFAAKKFAKALF 17

RESULT 19
US-09-820-053A-15
Sequence 15; Application US/09/820053A
Publication No. US20030084243A1
GENERAL INFORMATION:
1 APPLICANT: OWEN, DONALD R.
2 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES
3 FILE REFERENCE: HELIX027
4 CURRENT APPLICATION NUMBER: US/09/820,053A
5 CURRENT FILING DATE: 2001-03-28
6 NUMBER OF SEQ ID NOS: 165
7 SOFTWARE: PATENTIN Ver. 2.1
8 SEQ ID NO: 15
9 LENGTH: 24
10 TYPE: PRT
11 ORGANISM: ARTIFICIAL SEQUENCE
12 FEATURE:
13 OTHER INFORMATION: SYNTHETIC SEQUENCE
14 NAME/KEY: MOD_PES
15 LOCATION: (23)
16 OTHER INFORMATION: AMIDATION
17 US-09-820-053A-15

Query Match      63.68; Score 42; DB 11; Length 23;
Best Local Similarity 78.68; Pred. No. 2.9;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

CY 2 AKALAKALKALK 15
1 | | | | | | | | |
DB 2 AKALAKALKALK 15

RESULT 20
US-09-820-053A-158
Sequence 158; Application US/09/820053A
Publication No. US20030084243A1
GENERAL INFORMATION:
1 APPLICANT: OWEN, DONALD R.
2 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES
3 FILE REFERENCE: HELIX027
4 CURRENT APPLICATION NUMBER: US/09/820,053A
5 CURRENT FILING DATE: 2001-03-28
6 NUMBER OF SEQ ID NOS: 165
7 SOFTWARE: PATENTIN Ver. 2.1
8 SEQ ID NO: 158
9 LENGTH: 24
10 TYPE: PRT
11 ORGANISM: ARTIFICIAL SEQUENCE
12 FEATURE:
13 OTHER INFORMATION: SYNTHETIC SEQUENCE
14 NAME/KEY: MOD_PES
15 LOCATION: (23)
16 OTHER INFORMATION: AMIDATION
17 US-09-820-053A-158

Query Match      63.68; Score 42; DB 11; Length 24;
Best Local Similarity 78.68; Pred. No. 2.9;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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NAME/KEY: MOD_RES
 LOCATION: (15)
 OTHER INFORMATION: AMINATION
 US-09-820-053A-16

Query Match 62.1% Score 41: DB 11: Length 15:
 Best Local Similarity 74.8% Pred. No. 2.6:
 Matches 11: Conservative 0: Mismatches 4: Indels 0: Gaps 0:

QY 1 FAKAKAKAKAKAL 15
 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAKAL 15

RESULT 26
 US-10-109-171-22

Sequence 22, Application US/10109171
 Publication No. US20040109452A1

GENERAL INFORMATION:

APPLICANT: Owen, Donald R.

TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE

FILE REFERENCE: HELIX028

CURRENT APPLICATION NUMBER: US/10-109-171

CURRENT FILING DATE: 2002-03-28

NUMBER OF SEQ ID NOS: 165

SOFTWARE: Patent In Ver. 2.1

SEQ ID NO 22

LENGTH: 15

TYPE: PRT

ORGANISM: ARTIFICIAL SEQUENCE

FEATURE:

OTHER INFORMATION: SYNTHETIC SEQUENCE

NAME/KEY: MOD_RES

LOCATION: (15)

OTHER INFORMATION: AMINATION

US-10-109-171-22

Query Match 62.1% Score 41: DB 15: Length 15:
 Best Local Similarity 74.8% Pred. No. 2.6:
 Matches 11: Conservative 0: Mismatches 4: Indels 0: Gaps 0:

QY 1 FAKAKAKAKAKAL 15
 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAKAL 15

RESULT 27
 US-09-820-053A-16

Sequence 16, Application US/09820053A
 Publication No. US20040083241A1

GENERAL INFORMATION:

APPLICANT: Owen, Donald R.

TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES

FILE REFERENCE: HELIX027

CURRENT APPLICATION NUMBER: US/09-820-053A

CURRENT FILING DATE: 2001-03-28

NUMBER OF SEQ ID NOS: 165

SOFTWARE: Patent In Ver. 2.1

SEQ ID NO 16

LENGTH: 16

TYPE: PRT

ORGANISM: ARTIFICIAL SEQUENCE

FEATURE:

OTHER INFORMATION: SYNTHETIC SEQUENCE

NAME/KEY: MOD_RES

LOCATION: (16)

OTHER INFORMATION: AMINATION

US-09-820-053A-16

Query Match 62.1% Score 41: DB 11: Length 16:
 Best Local Similarity 76.9% Pred. No. 2.8:
 Matches 10: Conservative 0: Mismatches 2: Indels 0: Gaps 0:

QY 1 FAKAKAKAKAKAL 15
 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAKAL 15

QY 1 FAKAKAKAKAK 13
 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAK 13

RESULT 28

US-10-109-171-16
 Sequence 16, Application US/10109171
 Publication No. US20040109452A1

GENERAL INFORMATION:

APPLICANT: Owen, Donald R.

TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE

FILE REFERENCE: HELIX028

CURRENT APPLICATION NUMBER: US/10-109-171

CURRENT FILING DATE: 2002-03-28

NUMBER OF SEQ ID NOS: 165

SOFTWARE: Patent In Ver. 2.1

SEQ ID NO 16

LENGTH: 16

TYPE: PRT

ORGANISM: ARTIFICIAL SEQUENCE

FEATURE:

OTHER INFORMATION: SYNTHETIC SEQUENCE

NAME/KEY: MOD_RES

LOCATION: (16)

OTHER INFORMATION: AMINATION

US-10-109-171-16

Query Match 62.1% Score 41: DB 15: Length 16:
 Best Local Similarity 76.9% Pred. No. 2.8:
 Matches 10: Conservative 0: Mismatches 2: Indels 0: Gaps 0:

QY 1 FAKAKAKAKAK 13
 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAK 13

RESULT 29
 US-09-820-053A-161

Sequence 161, Application US/09820053A
 Publication No. US20040083241A1

GENERAL INFORMATION:

APPLICANT: Owen, Donald R.

TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES

FILE REFERENCE: HELIX027

CURRENT APPLICATION NUMBER: US/09-820-053A

CURRENT FILING DATE: 2001-03-28

NUMBER OF SEQ ID NOS: 165

SOFTWARE: Patent In Ver. 2.1

SEQ ID NO 161

LENGTH: 23

TYPE: PRT

ORGANISM: ARTIFICIAL SEQUENCE

FEATURE:

OTHER INFORMATION: SYNTHETIC SEQUENCE

NAME/KEY: MOD_RES

LOCATION: (23)

OTHER INFORMATION: AMINATION

US-09-820-053A-161

Query Match 62.1% Score 41: DB 11: Length 23:
 Best Local Similarity 76.9% Pred. No. 4.1:
 Matches 10: Conservative 0: Mismatches 2: Indels 0: Gaps 0:

QY 1 FAKAKAKAKAKAL 15
 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
 ID 1 FAKAKAKAKAKAL 15

RESULT 30
 US-10-109-171-161

Sequence 161, Application US/10109171
 Publication No. US20040109452A1

GENERAL INFORMATION:

APPLICANT: Owen, Donald R.

TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE

FILE REFERENCE: HELIX028

CURRENT APPLICATION NUMBER: US/10-109-171

CURRENT FILING DATE: 2002-03-28

NUMBER OF SEQ ID NOS: 165

SOFTWARE: Patent In Ver. 2.1

SEQ ID NO 161

LENGTH: 16

TYPE: PRT

ORGANISM: ARTIFICIAL SEQUENCE

FEATURE:

OTHER INFORMATION: SYNTHETIC SEQUENCE

NAME/KEY: MOD_RES

LOCATION: (16)

OTHER INFORMATION: AMINATION

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GENERAL INFORMATION:
  APPLICATION: US-09-820-053a-43.rapb
  FILE REFERENCE: HELIX02
  CURRENT ATTENTION NUMBER: 002709171
  CURRENT FILING DATE: 2002-04-28
  NUMBER OF SEQ ID NOS: 165
  SOFTWARE: PatGen V09.2.1
  TYPE: PRF
  ORGANISM: ARTIFICIAL SEQUENCE
  FEATURE:
  NAME/KEY: M11111
  LOCATION: (12)
  OTHER INFORMATION: AMINATION
US-10 109 171-106

Query Match
  Query: 171-106
  Subject: 171-106
  Matches: 109 Conservative 07 Mismatches 27 Indels 02 Gaps 02

RESULT 34
US-09 820 053a 8
1 Sequence 8: Application US/09820053a
2 Publication No: US20030088245A1
3 GENERAL INFORMATION:
4 APPLICATION: 0009, Isolated R.
5 FILE REFERENCE: HELIX02
6 CURRENT ATTENTION NUMBER: 002709820053a
7 CURRENT FILING DATE: 2001-04-28
8 NUMBER OF SEQ ID NOS: 165
9 SOFTWARE: PatGen V09.2.1
10 SEQ ID NO: 8
11 LENGTH: 24
12 TYPE: PRF
13 ORGANISM: ARTIFICIAL SEQUENCE
14 FEATURE:
15 NAME/KEY: M11111
16 LOCATION: (24)
17 OTHER INFORMATION: AMINATION
US-09 820 053a 8

Query Match
  Query: 171-106
  Subject: 171-106
  Matches: 109 Conservative 07 Mismatches 27 Indels 02 Gaps 02

RESULT 35
US-10 109 171-8
1 Sequence 8: Application US/10109171
2 Publication No: US20030109452A1
3 GENERAL INFORMATION:
4 APPLICATION: 0009, Isolated R.
5 FILE REFERENCE: HELIX02
6 CURRENT ATTENTION NUMBER: 002709109171
7 CURRENT FILING DATE: 2002-04-28
8 NUMBER OF SEQ ID NOS: 165
9 SOFTWARE: PatGen V09.2.1
10 SEQ ID NO: 8
11 LENGTH: 24
12 TYPE: PRF
13 ORGANISM: ARTIFICIAL SEQUENCE
14 FEATURE:
15 NAME/KEY: M11111
16 LOCATION: (24)
17 OTHER INFORMATION: AMINATION
US-10 109 171-8

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GENERAL INFORMATION:
  APPLICATION: US-09-820-053a-43.rapb
  FILE REFERENCE: HELIX02
  CURRENT ATTENTION NUMBER: 002709171
  CURRENT FILING DATE: 2002-04-28
  NUMBER OF SEQ ID NOS: 165
  SOFTWARE: PatGen V09.2.1
  TYPE: PRF
  ORGANISM: ARTIFICIAL SEQUENCE
  FEATURE:
  NAME/KEY: M11111
  LOCATION: (12)
  OTHER INFORMATION: AMINATION
US-10 109 171-106

Query Match
  Query: 171-106
  Subject: 171-106
  Matches: 109 Conservative 07 Mismatches 27 Indels 02 Gaps 02

RESULT 34
US-09 820 053a 8
1 Sequence 8: Application US/09820053a
2 Publication No: US20030088245A1
3 GENERAL INFORMATION:
4 APPLICATION: 0009, Isolated R.
5 FILE REFERENCE: HELIX02
6 CURRENT ATTENTION NUMBER: 002709820053a
7 CURRENT FILING DATE: 2001-04-28
8 NUMBER OF SEQ ID NOS: 165
9 SOFTWARE: PatGen V09.2.1
10 SEQ ID NO: 8
11 LENGTH: 24
12 TYPE: PRF
13 ORGANISM: ARTIFICIAL SEQUENCE
14 FEATURE:
15 NAME/KEY: M11111
16 LOCATION: (24)
17 OTHER INFORMATION: AMINATION
US-09 820 053a 8

Query Match
  Query: 171-106
  Subject: 171-106
  Matches: 109 Conservative 07 Mismatches 27 Indels 02 Gaps 02

RESULT 35
US-10 109 171-8
1 Sequence 8: Application US/10109171
2 Publication No: US20030109452A1
3 GENERAL INFORMATION:
4 APPLICATION: 0009, Isolated R.
5 FILE REFERENCE: HELIX02
6 CURRENT ATTENTION NUMBER: 002709109171
7 CURRENT FILING DATE: 2002-04-28
8 NUMBER OF SEQ ID NOS: 165
9 SOFTWARE: PatGen V09.2.1
10 SEQ ID NO: 8
11 LENGTH: 24
12 TYPE: PRF
13 ORGANISM: ARTIFICIAL SEQUENCE
14 FEATURE:
15 NAME/KEY: M11111
16 LOCATION: (24)
17 OTHER INFORMATION: AMINATION
US-10 109 171-8

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US-109-171-112
1 Sequence 116: Application US/0109171
2 Publication No. US20000109452A1
3 ORGANISM: INFORMATION:
4 APPLICANT: OZON, Donald R.
5 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE
6 FILE REFERENCE: HELIX28
7 CURRENT APPLICATION NUMBER: 09/07109, 171
8 CURRENT FILING DATE: 2002-03-28
9 NUMBER OF SEQ ID NOS: 165
10 SOFTWARE: PatentIn Ver. 2.1
11 SEQ ID NO: 112
12 LENGTH: 17
13 TYPE: PRT
14 ORGANISM: ARTIFICIAL SEQUENCE
15 FEATURE:
16 NAME/KEY: MOD_RES
17 LOCATION: (17)
18 OTHER INFORMATION: AMIDATION
US-109-171-112

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Query Match 56.1% Score 37: DB 15: Length 17:
Best Local Similarity 81.8% Pred. No. 12:
Matches 9: Conservative 0: Mismatches 2: Indels 0: Gaps 0:
CY 4 ALKALAKA 14
DB 4 ALKALAKA 14

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RESULT 43
US-109-171-112
1 Sequence 116: Application US/0109171
2 Publication No. US20000109452A1
3 GENERAL INFORMATION:
4 APPLICANT: OZON, Donald R.
5 TITLE OF INVENTION: SHORT BIOACTIVE PEPTIDES AND METHODS FOR THEIR USE
6 FILE REFERENCE: HELIX28
7 CURRENT APPLICATION NUMBER: 09/07109, 171
8 CURRENT FILING DATE: 2002-03-28
9 NUMBER OF SEQ ID NOS: 165
10 SOFTWARE: PatentIn Ver. 2.1
11 SEQ ID NO: 115
12 LENGTH: 17
13 TYPE: PRT
14 ORGANISM: ARTIFICIAL SEQUENCE
15 FEATURE:
16 NAME/KEY: MOD_RES
17 LOCATION: (17)
18 OTHER INFORMATION: AMIDATION
US-109-171-112

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Query Match 56.1% Score 37: DB 15: Length 17:
Best Local Similarity 81.8% Pred. No. 12:
Matches 9: Conservative 0: Mismatches 2: Indels 0: Gaps 0:

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US 4 KALKALKAK 14
DB 4 KALKALKAK 14
1 Sequence 12: Application US/09765614B
2 Publication No. US2002010215A1
3 GENERAL INFORMATION:
4 APPLICANT: Nycomed Imaging AS
5 TITLE OF INVENTION: Improvements in or relating to
6 TITLE OF INVENTION: diagnostic/therapeutic
7 TITLE OF INVENTION: agents
8 FILE REFERENCE: REF/61000006/054

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1 CURRENT APPLICATION NUMBER: US/09765614B
2 CURRENT FILING DATE: 2001-07-10
3 NUMBER OF SEQ ID NOS: 31
4 SOFTWARE: PatentIn Ver. 2.1
5 SEQ ID NO: 12
6 LENGTH: 18
7 TYPE: PRT
8 ORGANISM: Artificial Sequence
9 FEATURE:
10 OTHER INFORMATION: Description of Artificial
11 OTHER INFORMATION: Sequence:Endothelial
12 OTHER INFORMATION: cell binding lipopeptide
13 NAME/KEY: MOD_RES
14 LOCATION: (1)
15 OTHER INFORMATION: 2-n-hexadecylstearyl lys
16 NAME/KEY: MOD_RES
17 LOCATION: (18)
18 OTHER INFORMATION: AMIDATION
US-09-765-614B-12

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```

Query Match 56.1% Score 37: DB 10: Length 18:
Best Local Similarity 81.8% Pred. No. 13:
Matches 9: Conservative 0: Mismatches 2: Indels 0: Gaps 0:
CY 4 ALKALAKA 14
DB 4 ALKALAKA 14

```

```

RESULT 45
US-09-925-715-12
1 Sequence 12: Application US/09925715
2 Patent No. US20020102217A1
3 GENERAL INFORMATION:
4 APPLICANT: Nycomed Imaging AS
5 TITLE OF INVENTION: Improvements in or relating to diagnostic/therapeutic
6 FILE REFERENCE: REF/61000006/054
7 CURRENT APPLICATION NUMBER: 09/09925715
8 CURRENT FILING DATE: 2001-08-10
9 NUMBER OF SEQ ID NOS: 27
10 SOFTWARE: PatentIn Ver. 2.1
11 SEQ ID NO: 12
12 LENGTH: 18
13 TYPE: PRT
14 ORGANISM: Artificial Sequence
15 FEATURE:
16 OTHER INFORMATION: Description of Artificial Sequence:Synthetic
17 OTHER INFORMATION: endothelial cell binding lipopeptide
18 NAME/KEY: MOD_RES
19 LOCATION: (1)
20 OTHER INFORMATION: 2-n-hexadecylstearyl lysine
21 NAME/KEY: MOD_RES
22 LOCATION: (18)
23 OTHER INFORMATION: AMIDATION
US-09-925-715-12

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Query Match 56.1% Score 37: DB 10: Length 18:
Best Local Similarity 81.8% Pred. No. 13:
Matches 9: Conservative 0: Mismatches 2: Indels 0: Gaps 0:

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US 4 ALKALAKA 14
DB 4 ALKALAKA 14

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Search completed: August 21, 2003, 08:24:45
Job time : 55 secs

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The first part of the document discusses the importance of maintaining accurate records of all transactions. It emphasizes that proper record-keeping is essential for the transparency and accountability of the organization. The document then outlines the specific procedures for recording transactions, including the use of standardized forms and the requirement for double-checking entries. It also addresses the need for regular audits to ensure the integrity of the data. The second part of the document focuses on the financial aspects of the organization's operations. It details the budgeting process, from the initial planning stage to the final execution. It highlights the importance of staying within the allocated budget and provides strategies for managing any potential overruns. The document also discusses the reporting requirements for financial performance, including the preparation of monthly and annual reports. The final part of the document covers the administrative and legal aspects of the organization's activities. It outlines the necessary steps for obtaining permits and licenses, as well as the requirements for maintaining compliance with relevant regulations. It also discusses the importance of clear communication and documentation in all administrative matters. The document concludes by reiterating the commitment to transparency and accountability, and expresses confidence in the organization's ability to achieve its goals.